

LIMITED PHASE II ENVIRONMENTAL SITE ASSESSMENT

CITY OF HOUSTON 9003 NORTH MAIN STREET HOUSTON, TX ATC PROJECT NO. 73.17331.0074

June 30, 2006

Prepared by:

ATC Associates Inc. 3928 Bluebonnet Drive Stafford, Texas 77477 Telephone: (281) 240-0154

Facsimile: (281) 240-8909

Prepared for:

The City of Houston 900 Bagby, 2nd Floor Houston, TX 77002 Telephone: (713) 247-3232

Facsimile: (713)

Patrick Dworaczyk, P.G., CAPM Project Manager

Mark Smith, P.G., CAPM Senior Geologist

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EXECUTIVE SUMMARY

ATC Associates completed a limited Phase II Environmental Site Assessment (ESA) of the City of Houston property located at 9003 North Main Street, in Houston, Texas (herein referred to as the subject property). This report details the completion of the Limited Phase II ESA, in accordance with applicable ASTM standards and accepted environmental practices.

This Limited Phase II ESA conducted on June 21, 2006 identified the presence of BTEX and formaldehyde in the soil and groundwater. The constituent concentrations identified during the Phase II ESA were evaluated with respect to previous concentrations associated with the leaking petroleum storage tank (LPST) case and the TRRP Tier 1 groundwater protective concentration levels (PCLs). BTEX concentrations detected during this assessment were above TCEQ action levels for petroleum underground storage tank sites. These concentrations were below the previous concentrations detected at the site when closure was granted in 1997. The evaluation also identified formaldehyde to be below the TRRP Tier 1 groundwater PCL.

Three soil borings were advanced during the course of this Limited Phase II ESA. Soil samples were collected at the 12 – 14 ft below ground surface (bgs) interval in B-1, and the 20-22 ft bgs interval in B-2 and B-3. One soil samples from each boring was analyzed for BTEX by EPA Method 8021B, TPH by TX Method 1005, and Formaldehyde by EPA Method 8315. Analytical results for BTEX and TPH were below previous concentrations under which the site closed and formaldehyde was below the TRRP Tier 1 Residential PCLs.

The soil borings were converted to temporary monitoring wells. Groundwater samples were collected, submitted to Severn Trent Laboratories (STL) and analyzed for BTEX by EPA Method 8021B, TPH by TX Method 1005, and Formaldehyde by EPA Method 8315.

The concentrations identified during the Phase II ESA were evaluated with respect to previous concentrations associated with LPST case and the TRRP Tier 1 groundwater protective concentration levels (PCLs). BTEX was detected above TCEQ action levels but were below previous concentrations at the site. Formaldehyde concentrations were below the TRRP Tier 1 Residential PCLs.

Based on the data provided during the course of this Limited Phase II ESA, it appears that no further actions are recommended for the subject property at this time.

1.0 INTRODUCTION

1.1 Purpose and Scope

ATC Associates (ATC) was contracted by the City of Houston to conduct a Limited Phase II ESA of the subject property located at 9003 North Main in Houston, TX.

The purpose of the Limited Phase II Environmental Site Assessment (ESA) is to assess impact to the property from the LPST release associated with the site as well as reported historical use as a funeral home. The Limited Phase II ESA was performed to assist in confirming the nature and extent of contamination in the soil and groundwater of the above-referenced property.

The scope of the Limited Phase II Investigation included subsurface soil sampling, the installation of three temporary groundwater monitoring points, and groundwater sampling via the installation of three soil borings and temporary monitoring wells.

1.2 Background

The property is owned by the City of Houston, and was formerly used as an office and storage facility by the Street Maintenance Department. The site contains a one-story office building with an attached warehouse area in the eastern section of the property, a metal warehouse building at the north end of the property, and a canopy covered truck wash area on the southwestern portion of the site. The site also contains a concrete paved storage area formerly used for gravel, soil and other bulk materials used for street maintenance. The subject property was also identified in the TCEQ database as a leaking petroleum storage tank (LPST) site. A total of four USTs located at the property were reportedly removed in 1992.

2.0 PREVIOUS INVESTIGATIONS

2.1 Phase I Environmental Site Assessments

Weston conducted a Phase I ESA which identified the following environmental concerns:

- The subject property was identified in the TCEQ database as a leaking petroleum storage tank (LPST) site. A total of four USTs located at the property were reportedly removed in 1992. Following removal of the USTs, an assessment and groundwater monitoring activities were performed at the site. Groundwater was impacted but it was determined that there were no apparent threats or impacts to receptors. TCEQ issued final closure of the LPST case in February 1998.
- The office building present at the property was constructed prior to the 1970s and may contain asbestos and lead-based paint.
- The property was reportedly used as a funeral home prior to 1984; however, no documentation is available to confirm the former presence of a funeral home on the site.

2.2 Phase II Environmental Site Assessment

Carter and Burgess conducted Site Assessment activities at the site under the TCEQs LPST program in June 1995. Three groundwater monitoring wells were installed and soil and groundwater sampling was conducted. The soil and groundwater concentrations were above TCEQ action levels and the site was issued LPST number 104846. Groundwater monitoring was conducted and the site was closed in December 1997.

3.0 SOIL ASSESSMENT

3.1 Soil Boring Advancement and Sample Collection

On June 21, 2006, three soil borings were advanced with push-probe drilling equipment in the areas of the former USTs and near the back door of the office building. A site map depicting the soil boring location is included as Figure 1. The soil boring logs are provided in Appendix B.

During the advancement of the soil borings, soils were sampled continuously every 2 feet. One soil sample from each soil boring was selected for analysis; either the sample with the highest OVM reading or the sample at the soil groundwater interface was collected for laboratory analysis. Each soil sample was analyzed for benzene, toluene, ethylbenzene, and xylenes (BTEX) by EPA Method 8021B, total petroleum hydrocarbons (TPH) by Texas Method 1005, and formaldehyde by EPA Method 8315.

3.2 Laboratory Analysis

Analytical results of the soil samples collected by ATC indicated the presence of BTEX, and formaldehyde. Benzene concentrations ranged from below laboratory method detection limits in boring B-1 and B-3 to 0.072 mg/kg in B-1. Total BTEX concentrations ranged from below laboratory method detection limits in borings B-1 and B-3 to 2.982 mg/kg in B-2. TPH results were below laboratory method detection limits in the samples that were submitted for analysis. Formaldehyde was detected at concentrations of 2.520 mg/kg in B-1, 0.575 mg/kg in B-2, and 0.357 mg/kg in B-3. A copy of the laboratory analytical report by STL is provided in Appendix B.

The benzene concentration detected in B-2 is above the TCEQ action level for a UST site but is below the concentrations under which the site closed in 1997. The formaldehyde concentrations are below the TRRP Tier 1 Residential PCLS.

4.0 GROUNDWATER ASSESSMENT

4.1 Groundwater Sampling

The soil borings were converted to temporary groundwater monitoring wells. The temporary monitoring wells were constructed of threaded connection 1-inch ID, Schedule 40 PVC solid pipe, and 0.010-inch slotted PVC well screen.

On June 21, 2006, the three temporary monitoring wells were purged and sampled using a peristaltic pump. The wells were purged prior to sampling to reduce the amount of sediment present in the groundwater samples.

4.2 Laboratory Analysis

The analytical results of the groundwater samples obtained by ATC from the temporary monitoring wells on June 21, 2006 indicated concentrations of BTEX and formaldehyde. Benzene concentrations ranged from below laboratory method detection limits in B-2 and B-3 to 25.6 ug/L in B-1. Total BTEX concentrations ranged from below laboratory detection limits in B-3 to 561 ug/L in B-1. Formaldehyde concentrations were 27.6 ug/L in B-1, 46.4 ug/L in B-2, and 9.8 ug/L in B-3. TPH concentrations were below laboratory detection limits in the three borings. A copy of the laboratory analytical report is included in Appendix B

5.0 REGULATORY EVALUATION

Analytical laboratory results from this Limited Phase II ESA were compared to action levels established by the TCEQ and to applicable TRRP PCL. In addition, BTEX and TPH concentrations were compared to previous concentrations from when the site was in the TCEQ LPST program.

5.1 Soils

The benzene concentration detected in B-1 is above the TCEQ action levels for UST sites but is less than previous concentrations under which the site closed in 1997. The formaldehyde concentrations detected in the groundwater samples collected from the temporary monitoring wells was below the TRRP Tier I Residential PCLs.

5.2 Groundwater

The benzene concentration detected in B-1 is above the TCEQ action levels for UST sites but is less than previous concentrations under which the site closed in 1997. The formaldehyde concentrations detected in the groundwater samples collected from the temporary monitoring wells was below the TRRP Tier I Residential PCLs.

6.0 QUALITY CONTROL/QUALITY ASSURANCE

6.1 Decontamination Procedures

Drill operations were conducted using hydraulic direct push rig, with plastic sleeves, which were replaced after each 4-foot push. The auger drill rig used metal samplers which were cleaned in an alconox solution after each sample.

6.2 Field QA/QC Procedures

Soil samples submitted for laboratory analysis were contained within a Teflon-lined glass jar, placed on ice, and transported to the laboratory for analysis. Soil samples submitted to the laboratory were analyzed for BTEX, TPH and formaldehyde.

The groundwater samples submitted for laboratory analysis were contained within the appropriate containers, placed on ice, and transported to the laboratory for analysis. The groundwater sample submitted to the laboratory was analyzed for BTEX, TPH and formaldehyde.

Each sample was labeled and secured to preserve the integrity of the identification, from the time the sample was collected until it was opened at the laboratory. For each sample, the sample container label and chain-of-custody form were completed. Soil and groundwater samples were immediately placed in a cooler containing ice or frozen ice packs and hand delivered to the laboratory.

6.3 Sample Quality Control/Quality Assurance

Various QA/QC procedures were followed by the environmental laboratory. Prior to initiating analysis, it is required to establish that a given instrument meets the method tuning standard. The calibration of each instrument was verified at frequencies specified in the EPA approved methods. A new standard curve must be prepared as specified in each method per EPA Method SW-846.

Prior to analysis, instruments are required to be calibrated by the appropriate procedure. Each calibration standard was tabulated and the retention times recorded. The laboratory QA/QC results are provided in Appendix C.

7.0 CONCLUSIONS AND RECOMMENDATIONS

7.1 Conclusions

ATC has provided the following conclusions of this Phase II Site Assessment based on the field activities conducted on June 21, 2006 at the subject property, and on laboratory analytical data of media samples collected by ATC.

- The soils at the site consist of silty clays. Groundwater was encountered at a depth of approximately 15 to 21 feet bgs. Three temporary monitoring wells were installed to a maximum depth of 25 feet bgs.
- Analytical results of soil samples collected during the advancement of three soil borings along the west/southwest property boundary indicated the presence of BTEX and formaldehyde
- Formaldehyde concentrations in soil and groundwater were below the TRRP Tier I Residential PCLs

Project No. 73.27434.0032 ATC Associates Inc.

7.2 Recommendations

Based on the analytical results of soil and groundwater samples collected during this Limited Site Assessment, no further action is recommended at this time. Even though the Benzene concentrations were above TCEQ action levels, they were below the previous soil and groundwater concentrations detected at the site under which site closure was achieved.

TABLES

TABLE 1 SOIL ANALYTICAL RESULTS City of Houston 9003 North Main Houston, Texas

			Domoznod	Tologo	Cebalkonana	Joneph. A	Total BTEV	L	Fotal Petroleum Hydrocarbons	a Hydrocarbor	SL	
Well	Date	Date Depth (feet)	(mg/kg)	(mg/kg)	(mg/kg) (mg/kg) (mg/kg)	Ayienes (mg/kg)	(mg/kg)	C6-C12	i	C6-C28	Total	Formaldehyde
			(99)	/BB	(B., B.,)	ê .	(9	(mg/kg)	(mg/kg)	(mg/kg)	C6-C35	(mg/kg)
B-1	06/21/06	06/21/06 12-14	<0.0056	<0.00774	<0.00742	<0.0233	QN	<6.57	<13.8	<13.8	<13.8	2.520
B-2	B-2 06/21/06 20-22	20-22	0.0721	0.155	0.785	1.970	2.982	<6.32	<13.3	<13.3	<13.3	0.575
B-3	B-3 06/21/06	20-22	<0.00539	<0.00745	<0.00715	<0.0224	ND	<6.33	<13.3	<13.3	<13.3	0.357

Notes:

mg/kg - milligrams per kilogram or parts per million.

BTEX - benzene, toluene, ethylbenzene, and xylenes by EPA Method 8021B.

ND - not detected above laboratory detection limits.

NA - not available.

TABLE 2 GROUNDWATER ANALYTICAL RESULTS City of Houston 9003 North Main Houston, Texas

		Renzene	Toluono	T the lates	V.40	Tarana 1-1-T	L	Total Petroleum Hydrocarbons	Hydrocarbon	SI	
Well	Date	(ng/L)	(ug/L)	(ug/L)	(ug/L)	(ue/L)	C6-C12	C12-C28	C6-C28	Total	Formaldehyde
				,	, D		(mg/L)		(mg/L)	C6-C35	(ng/L)
B-1	06/21/06	25.6	7.1	131	396.9	561	<0.48	<0.83	<0.83	<0.83	27.6
B-2	90/17/90	<0.170	<0.190	0.67	2.37	3.40	<0.48	<0.83	<0.83	<0.83	46.4
B-3	06/21/06	<0.170	<0.190	<0.230	<0.560	QN	<0.48	<0.84	<0.84	<0.84	8.6

Notes:

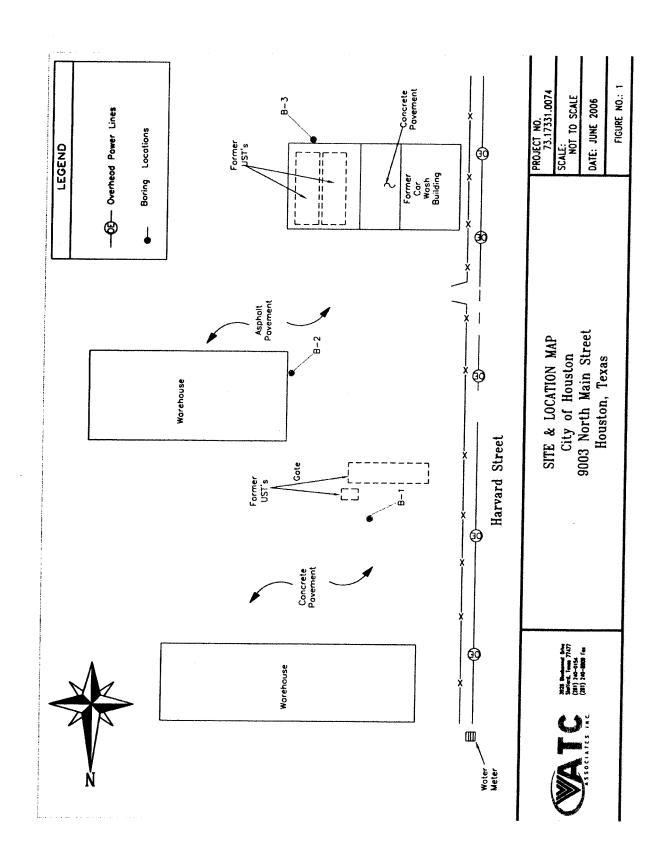
mg/kg - milligrams per kilogram or parts per million.

BTEX - benzene, toluene, ethylbenzene, and xylenes by EPA Method 8021B.

ND - not detected above laboratory detection limits.

NA - not available.

FIGURES



APPENDIX A **SOIL BORING LOGS**

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BAC	KFILL MATERIAL►										
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							2				
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50 — J											
POI	RING DESIGNATION			Γ					F	AGE	NUMBER

PROJECT► COH 9003 N. Main		Bori	na	PROJE	CT I	NUM	BER.	7 .	3.173	331.0074
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CHECKED BY>			9	СОМРІ	ETIC	ON D	ATE	>		
GROUND SURFACE ELEVATION DATUM (FT-MS	SL) ►		DRIL	ING C	ОМР	ANY	► Alp	ine		
DRILLING EQUIPMENT>			DRIL	ER►						
BORING DEPTH(FT)► \	WELL DEPTH(F	T) ►	WATE	R DEP	TH(F	T)-I	nitia	l:		Complete:
WELL MATERIALS ►			OVM,	OVA ►	OVM					
BACKFILL MATERIAL ►					······	,	,			
E LITH	IOLOGY				T		1	AMPL	E	-
O DE	RIPTION				GRAPHIC	OVM/OVA (PPM)	RECOVERY %	TIME	NUMBER	COMMENTS
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_										
						•				
Dark gray, silty clay Gray, silty clay— moist few Ca						0				
Gray, silty clay— moist few Ca	Nodules									
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Gray and brown clay— moist, fir	rm, Fe stains									
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30-										
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	CKED BY►	LU	<u>y</u>	СОМР	LETIC	ON D	DATE	>		
GRO ELE	UND SURFACE VATION DATUM (FT-MSL)►		DRIL	ING C	OMP	ANY	► Alp	ine		
	LING EQUIPMENT►		DRIL	_ER ►						
BOR	ING DEPTH(FT)► WELL DEPTH(F	-T) ►	WATE	R DEP	тн(ғ	T)-I	nitia	l:		Complete:
WEL	L MATERIALS ►		OVM,	OVA ►	OVM					
BAC	KFILL MATERIAL►									
(F	LITHOLOGY						i	AMPL	E.	
о рертн (DESCRIPTION				GRAPHIC	OVM/OVA (PPM)	RECOVERY %	TIME	NUMBER	COMMENTS
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						0				
10-	Fe staining below 10'									
						0				
	Red and gray clay, moist, stiff									
15						0				
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30-										
во	RING DESIGNATION B-3	ASSOC	T C		C.	*** ***********************************		F		NUMBER OF 1

APPENDIX B

LABORATORY REPORTS AND CHAIN-OF-CUSTODY DOCUMENTATION



ANALYTICAL REPORT

JOB NUMBER: 318054 Project ID: 9003 N. MAIN

Prepared For:

ATC Associates, Inc. 3928 Bluebonnet Drive Stafford, TX 77477

Attention: Patrick Dworaczyk

Date: 07/10/2006

Signature

Name: Dean A. Joiner

Title: Project Manager II

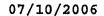
E-Mail: djoiner@stl-inc.com

Severn Trent Laboratories

6310 Rothway Drive Houston, TX 77040

PHONE: 713-690-4444

TOTAL NO. OF PAGES 48





Patrick Dworaczyk
ATC Associates, Inc.
3928 Bluebonnet Drive
Stafford, TX 77477

Reference:

Project : 9003 N. MAIN

Project No. : 318054
Date Received : 06/21/2006
STL Job : 318054

Dear Patrick Dworaczyk:

Enclosed are the analytical results for your project referenced above. The following samples are included in the report.

- 1. B-1 12-14
- 2. B-2 20-22
- 3. B-3 20-22
- 4. B-1
- 5. B-2
- 6. B-3
- 7. TRIP BLANK

All hold times were met for the tests performed on these samples.

Enclosed, please find the Quality Control Summary. All quality control results for the QC batch that are applicable to the sample(s) are acceptable except as noted in the QC batch reports.

The test results in this report meet all NELAP requirements for STL Houston's NELAP accredited parameters. Any exceptions to NELAP requirements will be noted and included in a case narrative as a part of this report.

If the report is acceptable, please approve the enclosed invoice and forward it for payment.

Thank you for selecting Severn-Trent Laboratories to serve as your analytical laboratory on this project. If you have any questions concerning these results, please feel free to contact me at any time.

We look forward to working with you on future projects.

Singerely,

Dean A. Joiner Project Manager



Table 1

Cross-Reference Field Sample Identifications and Laboratory Identifications

Field Identification	Laboratory Identification	8021B	TX1005	8315	Comment
B-1 12-14	318054-1	X	X	X	
B-2 20-22	318054-2	X	X	×	
B-3 20-22	318054-3	X	X	×	
B-1	318054-4	X	X	×	
B-2	318054-5	X	X	×	
B-3	318054-6	X	X	×	
TRIP BLANK	318054-7				Trip Blank; Not on C-O-C; No Tests Assigned

Appendix A Laboratory Data Package Cover Page

This data package consists of:

- This signature page, the laboratory review checklist, and the following reportable data:
- R1 Field chain-of-custody documentation;
- Sample identification cross-reference;
- R3 Test reports (analytical data sheets) for each environmental sample that includes:
 - a) Items consistent with NELAC 5.13 or ISO/IEC 17025 Section 5.10
 - b) dilution factors,
 - c) preparation methods,
 - d) cleanup methods, and
 - e) if required for the project, tentatively identified compounds (TICs).
- R4 Surrogate recovery data including:
 - a) Calculated recovery (%R), and
 - b) The laboratory's surrogate QC limits.
- Test reports/summary forms for blank samples; R5
- Test reports/summary forms for laboratory control samples (LCSs) including:
 - a) LCS spiking amounts,
 - b) Calculated %R for each analyte, and
 - c) The laboratory's LCS QC limits.
- Test reports for project matrix spike/matrix spike duplicates (MS/MSDs) including: R7
 - a) Samples associated with the MS/MSD clearly identified,
 - b) MS/MSD spiking amounts,
 - c) Concentration of each MS/MSD analyte measured in the parent and spiked samples,
 - d) Calculated %Rs and relative percent differences (RPDs), and
 - e) The laboratory's MS/MSD QC limits
- R8 Laboratory analytical duplicate (if applicable) recovery and precision:
 - a) the amount of analyte measured in the duplicate,
 - b) the calculated RPD, and
 - c) the laboratory's QC limits for analytical duplicates.
- R9 List of method quantitation limits (MQLs) for each analyte for each method and matrix;
- R10 Other problems or anomalies.
- The Exception Report for every "No" or "Not Reviewed (NR)" item in laboratory review checklist.

Release Statement: I am responsible for the release of this laboratory data package. This data package has been reviewed by the laboratory and is complete and technically compliant with the requirements of the methods used, except where noted by the laboratory in the attached exception reports. By me signature below, I affirm to the best of my knowledge, all problems/anomalies, observed by the laboratory as having the potential to affect the quality of the data, have been identified by the laboratory in the Laboratory Review Checklist, and no information or data have been knowingly withheld that would affect the quality of the data.

Check, if applica	able: [] This laborat	ory is an in-hous	se laboratory controlled by	y the p	person	la tha
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APAR) in w	hich these data are used	is responsible to	r releasing this data packs	rgc an	,	, 516,11414
affirming th	e above release statement	t ip true.				
	Washington	Nhor	Laboratory Director	7	112	100
Norman Flynn	Composit	0	Official Title (printed)		Date	L
Name (Printed)	Signature	()				

Lab	orato	ry Name: STL-Houston LRC Da	te: 06/26/06					
			ory Job Number: 318054					
<u> </u>			tch Number(s): 157296 (Soil)-TX10	05			~~~~	
			tell Number(s). 137290 (30h)-17(10h	Yes	No	NIA3	NR ⁴	ED+
# ¹	A ²	Description		105	INO	IVA	INK	5
		Chain-of-custody (C-O-C)					4,10	
RI	OI	Did samples meet the laboratory's standard conditions of sample a	cceptability upon receipt?		X			1
		Were all departures from standard conditions described in an excep		X				
R2	OI	Sample and quality control (QC) identification				7.2		
		Are all field sample ID numbers cross-referenced to the laboratory	ID numbers?	X				
	1	Are all laboratory ID numbers cross-referenced to the corresponding		X				
R3	OI	Test reports						
		Were all samples prepared and analyzed within holding times?		X				
		Other than those results < MQL, were all other raw values bracket	ed by calibration standards?	X				
		Were calculations checked by a peer or supervisor?		X				
ļ		Were all analyte identifications checked by a peer or supervisor?		X				
		Were sample quantitation limits reported for all analytes not detect	ed?	X				
		Were all results for soil and sediment samples reported on a dry we		X				
	ĺ	Were % moisture (or solids) reported for all soil and sediment sam		X				
		If required for the project, TICs reported?				X		
R4	o	Surrogate recovery data						
	 _ _ _ _ _ _ _ _ _ _ _ _ _ 	Were surrogates added prior to extraction?		X				
	1	Were surrogate percent recoveries in all samples within the laborate	ory OC limits?	X				\vdash
R5	OI	Test reports/summary forms for blank samples	ory QC mmo.	 				
-	101	Were appropriate type(s) of blanks analyzed?		X				\vdash
l		Were blanks analyzed at the appropriate frequency?		X				-
		Were method blanks taken through the entire analytical process, in	cluding prepa/ation and if	X				
		applicable, cleanup procedures?	cidding preparation and, it	1				
		Were blank concentrations < MQL?		X				一
R6	OI	Laboratory control samples (LCS):			 -			
100	<u> </u>	Were all COCs included in the LCS?		-	X			2
l	1	Was each LCS taken through the entire analytical procedure, inclu	ling area and cleanum stems?	X	1			F
		Were LCSs analyzed at the required frequency?	ing prep and eleurop steps:	X				
	1	Were LCS (and LCSD, if applicable) %Rs within the laboratory Q	? limite?	X				
	1	Does the detectability data document the laboratory's capability to		Λ		X		-
		to calculate the SQLs?	detect the COCs at the MDL used			^		
İ		Was the LCSD RPD within QC limits?		X	-			
R7	01	Matrix spike (MS) and matrix spike duplicate (MSD) data		^				
<u> </u>	101	Were the project/method specified analytes included in the MS and	MSD2	х				-
		Were MS/MSD analyzed at the appropriate frequency?	INOD:	X				
İ		Were MS (and MSD, if applicable) %Rs within the laboratory QC	limite?	<u>^</u>	-		X	3
1	l	Were MS/MSD RPDs within laboratory QC limits?	mms:	-			X	3
R8	OI	Analytical duplicate data			-			<u> </u>
No.	101	Were appropriate analytical duplicates analyzed for each matrix?				x		
		Were analytical duplicates analyzed at the appropriate frequency?				$\frac{\lambda}{X}$		-
			? limito?			$\hat{\mathbf{x}}$		-
-	-	Were RPDs or relative standard deviations within the laboratory Q	C mms;					
R9	OI	Method quantitation limits (MQLs):	inte makaga?	X				
1		Are the MQLs for each method analyte included in the laboratory		X				
}		Do the MQLs correspond to the concentration of the lowest non-zer	to campianon standaro?	X				-
D-10	<u></u>	Are unadjusted MQLs included in the laboratory data package?		^				
R10	OI	Other problems/anomalies	T DC and ED9	х			-	
l		Are all known problems/anomalies/special conditions noted in this		X				-
		Were all necessary corrective actions performed for the reported day. Was applicable and available technology used to lower the SQL to		X	-			-
		was applicable and available technology used to lower the SQL to affects on the sample results?	minimize me manta interference	^				L
		<u></u>						

SX

- Items identified by the letter "R" must be included in the laboratory data package submitted in the TRRP-required report(s). Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.
 O= organic analyses; I = inorganic analyses (and general chemistry, when applicable);
- NA = Not applicable;
- NR = Not reviewed;
- ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

ΑI

Ap	pen	dix A (cont'd): Laboratory Review Checklis	st: Reportable Data					
Lab	orato	ry Name: STL-Houston	LRC Date: 06/26/06					
Proj	ect N	Iame: 9003 N. MAIN	Laboratory Job Number: 318054					
Rev	iewe	r Name: MW	Prep Batch Number(s): 157296 (Soil)-TX	1005				
#1	A ²	Description		Yes	No	NA3	NR ⁴	ER#5
S1		Initial calibration (ICAL)		1111			25.00	
~		Were response factors and/or relative response factors for each	h analyte within OC limits?	X		†	†	
		Were percent RSDs or correlation coefficient criteria met?	<u> </u>	X		 		
		Was the number of standards recommended in the method use	ed for all analytes?	X				
		Were all points generated between the lowest and highest stan		X				
		Are ICAL data available for all instruments used?		X			T	
		Has the initial calibration curve been verified using an approp	riate second source standard?	X	 	 	 	1
S2	OI	Initial and continuing calibration verification (ICCV and		1	1.10	1		
-	<u>~</u> _	Was the CCV analyzed at the method-required frequency?		X			1	1
		Were percent differences for each analyte within the method-	required OC limits?	X	 		 	†
		Was the ICAL curve verified for each analyte?		$\frac{\hat{\mathbf{x}}}{\mathbf{x}}$		 	 	
		Was the absolute value of the analyte concentration in the inor	rganic CCB < MDL?	† ^	 	X	 	
S3	0	Mass spectral tuning:	PARALLE STATE STATE AND ADDRESS OF THE PARALLES AND ADDRES	1	100	 ``	1 . 1	1.74
22		Was the appropriate compound for the method used for tuning	79	 	 	X	1	1
		Were ion abundance data within the method-required QC limit		 	 	X	 	
S4	0	Internal standards (IS):	ω:	 	 	1	 	+
34	<u> </u>	Were IS area counts and retention times within the method-red	united OC limite?	├		X		
OE	OI.	Raw data (NELAC section 1 appendix A glossary, and section 1	tion 5 12 on ISO/ISC 17025 section	-	 	<u> </u>	 	
S5	<u>OI</u>	Were the raw data (for example, chromatograms, spectral data		Х		├	 	
				$\frac{\hat{x}}{x}$	├	 	 	
06		Were data associated with manual integrations flagged on the	raw datar	 ^-				
S6	<u>o</u>	Dual column confirmation	-1009	-		X		
- I		Did dual column confirmation results meet the method-require	ea QC?			A	17.57	-
S7	<u>o_</u>	Tentatively identified compounds (TICs):	1:					
		If TICs were requested, were the mass spectra and TIC data st	ibject to appropriate checks?	 	11 2 1	X	1 1 2 3 3 3 1 1	
S8	1	Interference Check Sample (ICS) results:				-		1
		Were percent recoveries within method QC limits?		-	7.7.	X		-
S9	<u> </u>	Serial dilutions, post digestion spikes, and method of stand	ard additions				1 2 2 2	
		Were percent differences, recoveries, and the linearity within	the QC limits specified in the method?	 	ļ	X		
S10	<u>01</u>	Method detection limit (MDL) studies		ļ				
		Was a MDL study performed for each reported analyte?		X	 	ļ	<u> </u>	
		Is the MDL either adjusted or supported by the analysis of DC	Ss?	X		ļ	ļ	
S11	<u>OI</u>	Proficiency test reports:		<u> </u>	<u> </u>	 	<u> </u>	ļ
		Was the laboratory's performance acceptable on the applicable	e proficiency tests or evaluation studies?	X	<u> </u>			
S12	OI	Standards documentation			ļ	<u> </u>		1
		Are all standards used in the analyses NIST-traceable or obtain	ned from other appropriate sources?	X	<u> </u>			<u> </u>
S13	OI	Compound/analyte identification procedures						
		Are the procedures for compound/analyte identification documents	nented?	X	<u> </u>			<u> </u>
S14	OI	Demonstration of analyst competency (DOC)					1. 1.50	
		Was DOC conducted consistent with NELAC Chapter 5C or I		X				<u> </u>
j	-	Is documentation of the analyst's competency up-to-date and	on file?	X				
S15	OI	Verification/validation documentation for methods (NELA	C Chap 5 or ISO/IEC 17025 Section 5)	17.00	35, 3	7,45%		13174
		Are all the methods used to generate the data documented, ver	ified, and validated, where applicable?	X				
	~~	Laboratory standard operating procedures (SOPs):					1700	10.000
S16	OI							

Items identified by the letter "R" should be included in the laboratory data package submitted to the TCEQ in the TRRP-required report(s). Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.

O = organic analyses; I = inorganic analyses (and general chemistry, when applicable).

5

NA = Not applicable.

NR = Not Reviewed.

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

Appe	ndix A (cont'd): Laboratory Review Ch	ecklist: Exception Reports				
Labora	itory Name: STL-Houston	LRC Date: 06/26/06				
Project	Name: 9003 N. MAIN	Laboratory Job Number: 318054				
Reviev	ver Name: MW	Prep Batch Number(s): 157296 (Soil)-TX1005				
ER#	DESCRIPTION					
1	The temperature of the cooler received by the laboratory on 06/21/06 was above the acceptable range of 2.0-6.0 °C.					
2	The final concentration of any hydrocarbons d	required by the method, this range was not spiked into the LCS/LCSD. letected in this range was calculated from the response factor of the C12-raction efficiency of the C28-C35 range hydrocarbons was determined ns.				
3	The laboratory selected another client's sample	e to perform as the MS/MSD.				

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked on the LRC)

·		ry Name: STL-Houston LRC D	ate: 06/26/06				-	
			tory Job Number: 318054					
		· · · · · · · · · · · · · · · · · · ·				····		
Rev	iewer	r Name: MW Prep B	atch Number(s): 157449 (Water)-TXI					
# ¹	A ²	Description		Yes	No	NA ³	NR ⁴	ER#
		Chain-of-custody (C-O-C)			A			110011
R1	OI		cceptability upon receipt?	 	X			1
		Were all departures from standard conditions described in an exce		X				
R2	OI	Sample and quality control (QC) identification		-				
	-	Are all field sample ID numbers cross-referenced to the laboratory	ID numbers?	X				
	1	Are all laboratory ID numbers cross-referenced to the correspondi		X				
R3	OI	Test reports		 		-		
	 	Were all samples prepared and analyzed within holding times?		X				
		Other than those results < MQL, were all other raw values bracket	ed by calibration standards?	$\frac{\Lambda}{X}$				
	1	Were calculations checked by a peer or supervisor?	od by cantraction standards:	X	 			
		Were all analyte identifications checked by a peer or supervisor?		$\frac{\Lambda}{X}$				
		Were sample quantitation limits reported for all analytes not detec	ted?	X				
		Were all results for soil and sediment samples reported on a dry w			-	x		├
		Were % moisture (or solids) reported for all soil and sediment sam				X		
l	1		pies:			X		
R4	0	If required for the project, TICs reported?						
K4	<u> </u>	Surrogate recovery data						
		Were surrogates added prior to extraction?	001, 70	X				
200		Were surrogate percent recoveries in all samples within the labora	tory QC limits?	X				ļ
R5	OI	Test reports/summary forms for blank samples			ļ			
		Were appropriate type(s) of blanks analyzed?		X				ļ
		Were blanks analyzed at the appropriate frequency?		X	ļ			
		Were method blanks taken through the entire analytical process, in	icluding prepa/ation and, if	X				İ
l		applicable, cleanup procedures?		-,,				ļ
		Were blank concentrations < MQL?		X		-		 -
R6	OI	Laboratory control samples (LCS):						
		Were all COCs included in the LCS?			Х			2
	1	Was each LCS taken through the entire analytical procedure, inclu	ding prep and cleanup steps?	X				ļ
		Were LCSs analyzed at the required frequency?		X				ļ
		Were LCS (and LCSD, if applicable) %Rs within the laboratory Q		X				ļ
		Does the detectability data document the laboratory's capability to	detect the COCs at the MDL used			X		ĺ
		to calculate the SQLs?						ļ
		Was the LCSD RPD within QC limits?		X				<u> </u>
R7	OI	Matrix spike (MS) and matrix spike duplicate (MSD) data						ļ
		Were the project/method specified analytes included in the MS an	d MSD?	X				ļ
		Were MS/MSD analyzed at the appropriate frequency?		X				ļ
		Were MS (and MSD, if applicable) %Rs within the laboratory QC	limits?	X				ļ
	<u> </u>	Were MS/MSD RPDs within laboratory QC limits?		X				
R8	OI	Analytical duplicate data						ļ
		Were appropriate analytical duplicates analyzed for each matrix?				X		
		Were analytical duplicates analyzed at the appropriate frequency?				X		
		Were RPDs or relative standard deviations within the laboratory Q	C limits?			Х		Ĺ
R9	OI	Method quantitation limits (MQLs):						
		Are the MQLs for each method analyte included in the laboratory		X				
		Do the MQLs correspond to the concentration of the lowest non-ze	ero calibration standard?	X				
		Are unadjusted MQLs included in the laboratory data package?		Х				
R10	OI	Other problems/anomalies						
		Are all known problems/anomalies/special conditions noted in this	LRC and ER?	Х				
	1	Were all necessary corrective actions performed for the reported d	ata?	X				
		Was applicable and available technology used to lower the SQL to		X				
	l	affects on the sample results?						

SX

- Items identified by the letter "R" must be included in the laboratory data package submitted in the TRRP-required report(s). Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.
 O= organic analyses; I = inorganic analyses (and general chemistry, when applicable);
- NA = Not applicable; NR = Not reviewed;
- ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

Αl

Reviewer Name	Name: STL-Houston	LRC Date: 06/26/06					
#	e: 9003 N. MAIN	Laboratory Job Number: 318054					
# A Desc S1 OI Initia	me: MW	Prep Batch Number(s): 157449 (Water)-7	'X10)5	····	*******	
S1 OI Initial Were Was Were Was Was Was Was Was Was Were Was Were Were Were Were Were Were S6 O Dual Did d S7 O Tental Tental Tental Tental Tental Tental Tental Were S9 I Seria Were S9 I Seria Were S10 OI Meth Was Is the S11 OI Profi Was Is the S12 OI Stand Are a S13 OI Comp Are the S14 OI Demot S15 OI Verif S15 OI OI OI OI OI OI OI O			Yes		NA ³	NR ⁴	ER#
Were Were Was Were	tial calibration (ICAL)		103	110	INA	HAX.	Lin
Were Was Were	ere response factors and/or relative response factors for each	h analyta within OC limita?	X		 		
Was Were	ere percent RSDs or correlation coefficient criteria met?	n analyte within QC hmits?	X		 		
Were Are I	as the number of standards recommended in the method use	ad for all analytes?	X				
Are I	ere all points generated between the lowest and highest star		X		 	├	
Has t S2 OI Initia Was Was Was Was Was Was Were Were Were Were Were S5 OI Raw Were Were S6 O Dual Did d S7 O Tenta If TIC S8 I Inter Were S9 I Seria Were S11 OI Meth Was a Is the S12 OI Stanc Are a S13 OI Comp Are till S14 OI Demot Was I S6 OI Verif Was I S15 OI Verif Was I S15 OI Verif	e ICAL data available for all instruments used?	idard used to carculate the curve?	X		-	 	
S2 OI Initia Was Were Was Were	s the initial calibration curve been verified using an approp	riote second source standard?	X		 		
Was Were Was Were Was Were	tial and continuing calibration verification (ICCV and (^	66.1			
Were Was	is the CCV analyzed at the method-required frequency?	CCV) and continuing candiation	X				-
Was Was	ere percent differences for each analyte within the method-	required OC limite?	X				
Was Was	is the ICAL curve verified for each analyte?	required QC minits:	X		 		
S3 O Mass Were	is the real curve verified for each analyte? Is the absolute value of the analyte concentration in the ino	roanic CCR < MDI 2	_^_		X	-	
Was Were	is the absolute value of the analyte concentration in the mo	Iganic CCB < WDL?			^	10.00	35.5
Were	is the appropriate compound for the method used for tuning				X		
S4 O Inter	ere ion abundance data within the method-required QC limi				X		
Were Were							
S5 OI Raw Were	Were IS area counts and retention times within the method-required QC limits?		ļ		X		
Were Were	w data (NELAC section 1 appendix A glossary, and sec						
Were	ere the raw data (for example, chromatograms, spectral data		X			-	
S6 O Dual			X				
Did d S7 O Tents If TiC S8 I Inter Were S9 I Seria Were S10 OI Meth Was a Is the S11 OI Profit Was 1 S12 OI Stance Are a S13 OI Com Are tit S14 OI Demo Was 1 Is doc S15 OI Verif	Were data associated with manual integrations flagged on the raw data? Dual column confirmation		-		, 973,		
S7 O Tentic	I dual column confirmation results meet the method-require	MOC2			X		
If TIC S8 I Inter Were S9 I Seria Were S10 OI Meth Was a Is the S11 OI Profit Was 1 S12 OI Stance Are a S13 OI Com Are the S14 OI Demo Was 1 Is doc S15 OI Verif		ed QC;					
S8 I Inter	ntatively identified compounds (TICs): TICs were requested, were the mass spectra and TIC data so	which to appropriate shocked			X	31/25/10	
Were Were	erference Check Sample (ICS) results:	doject to appropriate checks?	9 (%)	in iga.	-^-	0.135.5	111
Seria Seria	errefrence Check Sample (1CS) results: ere percent recoveries within method QC limits?				Х		
Were Were			GgSA.				10,00
S10 OI Meth Was a Is the	rial dilutions, post digestion spikes, and method of stand are percent differences, recoveries, and the linearity within				Х		
Was a Is the	thod detection limit (MDL) studies	the QC mints specified in the method?	72.5	71.75	_^_		
Is the	is a MDL study performed for each reported analyte?		х				
S11 OI	he MDL either adjusted or supported by the analysis of DC	200	$\frac{\lambda}{X}$				
Was t S12 OI Stand Are a S13 OI Com Are tt S14 OI Demo Was l Is doc	oficiency test reports:	, JS :	^		1 77 1		
S12 OI Stand Are a	is the laboratory's performance acceptable on the applicable	e proficiency tests or evaluation studies?	Х				
Are a	ndards documentation	e protected y tests of evaluation studies?	<u>^</u>	77.7		1 12 1	
S13 OI Com Are th S14 OI Demo Was I Is doc S15 OI Verif	all standards used in the analyses NIST-traceable or obtain	ned from other engraprists courses?	Х				
Are the S14 OI Demo Was I Is doc S15 OI Verif	mpound/analyte identification procedures	ned from other appropriate sources:	^	75.7	5.7 (4	1,75,175	Jan 12, 1
Name	the procedures for compound/analyte identification documents	nantad?	X				
Was I Is doc S15 OI Verif		AVIIIO I	7			3355	: दिवसी र
Is doc S15 OI Verif	monstration of analyst competency (DOC) s DOC conducted consistent with NELAC Chapter 5C or I	SO/IEC 42	\mathbf{x}				
S15 OI Verif	s DOC conducted consistent with NEEAC Chapter 3C of I		$\frac{\hat{\mathbf{x}}}{\mathbf{x}}$				
AT A CLI	rification/validation documentation for methods (NELA			199	127	4,55	VA.
Aran	e all the methods used to generate the data documented, ver	rified and validated where applicable?	Х				V17411
		ined, and vanuated, where applicable?					7 - 4
S16 OI Labo	boratory standard operating procedures (SOPs): aboratory SOPs current and on file for each method performs.		x				

Items identified by the letter "R" should be included in the laboratory data package submitted to the TCEQ in the TRRP-required report(s).

Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.

O = organic analyses; I = inorganic analyses (and general chemistry, when applicable).

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NR = Not Reviewed.

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

Appendix A (cont'd): Laboratory Review Checklist: Exception Reports						
Labora	tory Name: STL-Houston	LRC Date: 06/26/06				
Project Name: 9003 N. MAIN		Laboratory Job Number: 318054				
Reviewer Name: MW		Prep Batch Number(s): 157449 (Water)-TX1005				
ER#1	DESCRIPTION					
1	The temperature of the cooler received by the	laboratory on 06/21/06 was above the acceptable range of 2.0-6.0 °C.				
2	Since calibration to the C28-C35 range is not required by the method, this range was not spiked into the LCS/LCSD. The final concentration of any hydrocarbons detected in this range was calculated from the response factor of the C12-					
	C28 hydrocarbons. Based on this fact, the extraction efficiency of the C28-C35 range hydrocarbons was determined from the recovery of the C12-C28 hydrocarbons.					

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked on the LRC)

Lab	orato	ry Name: STL-Houston LR	C Date: 06/26/06					
			poratory Job Number: 318054					
<u> </u>				r				
#I			p Batch Number(s): 157581 (Soil)-BTEX		λī-	NA ³	NID 4	TOTAL!
#	A	Description Co. Co. Co. Co. Co. Co. Co. Co. Co. Co.		Yes	No	INA	NR⁴	ER#
nı	01	Chain-of-custody (C-O-C)				59,5		(core)
R1	U	Did samples meet the laboratory's standard conditions of samp		 	X	ļ		1
	-	Were all departures from standard conditions described in an e	exception report?	X	ļ			
R2	OI	Sample and quality control (QC) identification						111
		Are all field sample ID numbers cross-referenced to the labora		X		ļ		
72.0	-	Are all laboratory ID numbers cross-referenced to the correspondence	onding QC data?	X				
R3	OI	Test reports				ļ		
		Were all samples prepared and analyzed within holding times		X	ļ	ļ		ļ
		Other than those results < MQL, were all other raw values bra-	cketed by calibration standards?	X		ļ		ļ
		Were calculations checked by a peer or supervisor?		X				
		Were all analyte identifications checked by a peer or supervisor		X	ļ	ļ		
		Were sample quantitation limits reported for all analytes not d		X			 	
		Were all results for soil and sediment samples reported on a dr		X				
	1	Were % moisture (or solids) reported for all soil and sediment	samples?	X				
	<u> </u>	If required for the project, TICs reported?				X		
R4	0	Surrogate recovery data						
		Were surrogates added prior to extraction?		X				
	<u> </u>	Were surrogate percent recoveries in all samples within the lab	oratory QC limits?		X			2
R5	OI	Test reports/summary forms for blank samples						
		Were appropriate type(s) of blanks analyzed?		X				
		Were blanks analyzed at the appropriate frequency?		X				
		Were method blanks taken through the entire analytical proces	s, including preparation and, if	X				
		applicable, cleanup procedures?						
		Were blank concentrations < MQL?		X				
R6	OI	Laboratory control samples (LCS):						
		Were all COCs included in the LCS?		X				
		Was each LCS taken through the entire analytical procedure, in	ncluding prep and cleanup steps?	X				
		Were LCSs analyzed at the required frequency?		X				~~~~
		Were LCS (and LCSD, if applicable) %Rs within the laborator	y QC limits?	X				
		Does the detectability data document the laboratory's capabilit	y to detect the COCs at the MDL used			X		
		to calculate the SQLs?						
		Was the LCSD RPD within QC limits?				X		
R7	OI	Matrix spike (MS) and matrix spike duplicate (MSD) data						
		Were the project/method specified analytes included in the MS	and MSD?	X				***************************************
		Were MS/MSD analyzed at the appropriate frequency?		X				
		Were MS (and MSD, if applicable) %Rs within the laboratory	OC limits?	х				
		Were MS/MSD RPDs within laboratory QC limits?	·	Х	$\neg \neg$			
R8	OI	Analytical duplicate data						
		Were appropriate analytical duplicates analyzed for each matri	x?			Х		
		Were analytical duplicates analyzed at the appropriate frequen				X		
		Were RPDs or relative standard deviations within the laborator				X		
R9	OI	Method quantitation limits (MQLs):						
	<u> </u>	Are the MQLs for each method analyte included in the laborate	orv data package?	X		1	\neg	
		Do the MQLs correspond to the concentration of the lowest no		$\frac{x}{x}$				
		Are unadjusted MQLs included in the laboratory data package		$\frac{\Lambda}{X}$				
R10	OI	Other problems/anomalies						
	"	Are all known problems/anomalies/special conditions noted in	this I RC and FR?	X				
		Were all necessary corrective actions performed for the reporte		$\frac{\hat{X}}{X}$				
		Was applicable and available technology used to lower the SQ		$\frac{\Delta}{X}$				
		affects on the sample results?	a to minimize the matrix interference	^	ļ		1	
	Ļ	Items identified by the letter "P" must be included in the laboratory data	The state of the s	لببا			!	

^{1.} Items identified by the letter "R" must be included in the laboratory data package submitted in the TRRP-required report(s). Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.

2. = organic analyses; I = inorganic analyses (and general chemistry, when applicable);

3. NA = Not applicable;

4. NR = Not reviewed;

5. ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

Lab	orato	ry Name: STL-Houston LR	C Date: 06/26/06					
Proj	ect N	lame: 9003 N. MAIN La	boratory Job Number: 318054					
Rev	iewei	r Name: ERA	ep Batch Number(s): 157581 (Soil)-BTF	X				,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
#1		Description		Yes	No	NA ³	NR ⁴	ER#
					77.5	9 11 1	77.5	
<u>S1</u>	OI	Initial calibration (ICAL) Were response factors and/or relative response factors for each	analyta within OC limits?	X			 	
		Were percent RSDs or correlation coefficient criteria met?	anaryte within OC mints:	X		 	┧	
		Was the number of standards recommended in the method used	for all analytee?	X		 	 	
		Were all points generated between the lowest and highest stand	lord used to calculate the curve?	X			 	
			lard used to calculate the curve:	X	-	 	 	\vdash
		Are ICAL data available for all instruments used? Has the initial calibration curve been verified using an appropri	isto second source standard?	X	 	 	 	
~	2.0	Initial and continuing calibration verification (ICCV and C	CV) and continuing calibration	<u> </u>				
S2	OI		CV) and continuing campi atton	X		 	 	\vdash
		Was the CCV analyzed at the method-required frequency? Were percent differences for each analyte within the method-re	equired OC limite?	X	\vdash	 	 	t
		were percent differences for each analyte within the method-re	Admed AC minust	$\frac{\Lambda}{X}$	 	 	 	
		Was the ICAL curve verified for each analyte? Was the absolute value of the analyte concentration in the inorg	ronio CCR < MDI 2	<u> </u>	 	X	 	1-
~~			gaine CCB < MDL!	20127	-	1-	-	
S3	0	Mass spectral tuning:	3		-	X	 	
		Was the appropriate compound for the method used for tuning		-		X	 	
~ -		Were ion abundance data within the method-required QC limits	S?		-	<u> </u>	 	1
<u>54</u>	0	O Internal standards (IS): Were IS area counts and retention times within the method-required QC limits?		X				┼──
		Were IS area counts and retention times within the method-red	f 12 YCO/TEC 17025		-	-	 	┼
S5_	OI	Raw data (NELAC section 1 appendix A glossary, and section	on 5.12 or 180/IEC 1/025 section	X			-	┼─
		Were the raw data (for example, chromatograms, spectral data)		X		 	┼	┼
		Were data associated with manual integrations flagged on the r	aw data?	<u> </u>	-	-	 	┼──
S6	0	Dual column confirmation	1000	X			-	┼
		Did dual column confirmation results meet the method-required	d QC?	1		 	1 222	┼─
<u>S7</u>	0	Tentatively identified compounds (TICs):				37	-	-
		If TICs were requested, were the mass spectra and TIC data su	bject to appropriate checks?		1, 1	X	-	+
S8	I	Interference Check Sample (ICS) results:		71, 11		X		-
		Were percent recoveries within method QC limits?			1000		1	+
<u>59</u>	I	Serial dilutions, post digestion spikes, and method of standa	ard additions			47	-	
		Were percent differences, recoveries, and the linearity within the	he QC limits specified in the method?		11.	X	 	┼
S10	OI	Method detection limit (MDL) studies		-		 	-	┼—
	<u> </u>	Was a MDL study performed for each reported analyte?		X		 	 	┼
		Is the MDL either adjusted or supported by the analysis of DC	Ss?	X		 	-	
S11	OI	Proficiency test reports:		125			 	┼
		Was the laboratory's performance acceptable on the applicable	proficiency tests or evaluation studies?	X		ļ	 	┼
S12	OI	Standards documentation		-		ļ	 	-
		Are all standards used in the analyses NIST-traceable or obtain	ed from other appropriate sources?	X		ļ	 	
S13	OI	Compound/analyte identification procedures				<u> </u>	-	-
		Are the procedures for compound/analyte identification docum	ented?	X	ļ	-	 	╀
S14	OI	Demonstration of analyst competency (DOC)				ļ	1-	4
		Was DOC conducted consistent with NELAC Chapter 5C or IS	SO/IEC 4?	X	 	<u> </u>	 	₩
	<u></u>	Is documentation of the analyst's competency up-to-date and o	n file?	X		ļ		—
S15	OI	Verification/validation documentation for methods (NELAC	C Chap 5 or ISO/IEC 17025 Section 5)	35.1				1
	1	Are all the methods used to generate the data documented, veri	fied, and validated, where applicable?	X				
S16	OI	Laboratory standard operating procedures (SOPs):					1 1 1 1 1	
	 ~~	Are laboratory SOPs current and on file for each method perform	rmed?	X			1	

Items identified by the letter "R" should be included in the laboratory data package submitted to the TCEQ in the TRRP-required report(s). Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.

O = organic analyses; I = inorganic analyses (and general chemistry, when applicable).

NA = Not applicable.

NR = Not Reviewed.

³

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

Apper	ndix A (cont'd): Laboratory Revie	w Checklist: Exception Reports				
Laborat	ory Name: STL-Houston	LRC Date: 06/26/06				
Project	Name: 9003 N. MAIN	Laboratory Job Number: 318054				
Review	er Name: ERA	Prep Batch Number(s): 157581 (Soil)-BTEX				
ER#1	DESCRIPTION					
1	The temperature of the cooler received	The temperature of the cooler received by the laboratory on 06/21/06 was above the acceptable range of 2.0-6.0 °C.				
2		The a,a,a-trifluorotoluene surrogate recoveries on both columns and the bromofluorobenzene recovery on column SPB-624 in sample 318054-2 were outside acceptance limits due to matrix interference.				

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked on the LRC)

Lab	orato	ry Name: STL-Houston LR	RC Date: 06/26/06					
		· · · · · · · · · · · · · · · · · · ·	boratory Job Number: 318054					
				T237				
			ep Batch Number(s): 157626 (Water)-BT			120.3	T A	
# ¹	A.	Description		Yes	No	NA ³	NR ⁴	ER#5
	~-	Chain-of-custody (C-O-C)						315 7
R1	OI	Did samples meet the laboratory's standard conditions of sam	ple acceptability upon receipt?	ļ	X	<u> </u>		1
	ļ	Were all departures from standard conditions described in an	exception report?	X				
R2	OI	Sample and quality control (QC) identification		<u> </u>				
		Are all field sample ID numbers cross-referenced to the labor		X				
	-	Are all laboratory ID numbers cross-referenced to the corresp	onding QC data?	X		<u> </u>	<u> </u>	
R3	OI	Test reports		 				
		Were all samples prepared and analyzed within holding times		X	<u> </u>			
		Other than those results < MQL, were all other raw values bra	acketed by calibration standards?	X	 	ļ		ļ
	İ	Were calculations checked by a peer or supervisor?		X		ļ		
		Were all analyte identifications checked by a peer or supervis		X	<u> </u>	ļ		
		Were sample quantitation limits reported for all analytes not of		X		ļ		
		Were all results for soil and sediment samples reported on a d		X		<u> </u>		
		Were % moisture (or solids) reported for all soil and sediment	t samples?	X		L		
	<u> </u>	If required for the project, TICs reported?				X		
R4	0	Surrogate recovery data						
		Were surrogates added prior to extraction?		X				
		Were surrogate percent recoveries in all samples within the la	boratory QC limits?		X			2
R5	OI	Test reports/summary forms for blank samples						
		Were appropriate type(s) of blanks analyzed?		X				
		Were blanks analyzed at the appropriate frequency?		X				
		Were method blanks taken through the entire analytical proce-	ss, including preparation and, if	X				
		applicable, cleanup procedures?						
		Were blank concentrations < MQL?		X				
R6	OI	Laboratory control samples (LCS):						
		Were all COCs included in the LCS?		X				
		Was each LCS taken through the entire analytical procedure,	including prep and cleanup steps?	X				
		Were LCSs analyzed at the required frequency?		X				
		Were LCS (and LCSD, if applicable) %Rs within the laborato		X				
		Does the detectability data document the laboratory's capabili	ty to detect the COCs at the MDL used			X		
		to calculate the SQLs?		\sqcup				
	~=	Was the LCSD RPD within QC limits?				X		
R7	OI	Matrix spike (MS) and matrix spike duplicate (MSD) data						
		Were the project/method specified analytes included in the M	S and MSD?	X				
		Were MS/MSD analyzed at the appropriate frequency?		X				
		Were MS (and MSD, if applicable) %Rs within the laboratory	QC limits?				X	3
		Were MS/MSD RPDs within laboratory QC limits?		 			X	3
R8	OI	Analytical duplicate data						
		Were appropriate analytical duplicates analyzed for each matr				X		
		Were analytical duplicates analyzed at the appropriate frequen	· · · · · · · · · · · · · · · · · · ·	lacksquare		X		
		Were RPDs or relative standard deviations within the laborato	ry QC limits?			X		
R9	<u>OI</u>	Method quantitation limits (MQLs):	MMMV - Magnetin (1975)	ليا				
ļ		Are the MQLs for each method analyte included in the laborat		X			$-\!\!\perp$	
		Do the MQLs correspond to the concentration of the lowest no		X				
		Are unadjusted MQLs included in the laboratory data package	37	X				
R10	OI	Other problems/anomalies		لييا				
		Are all known problems/anomalies/special conditions noted in		X				
		Were all necessary corrective actions performed for the report		X				
		Was applicable and available technology used to lower the SQ	L to minimize the matrix interference	X	- 1		1	
		affects on the sample results?						

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4. NR = Not reviewed;

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A	per	ndix A (cont'd): Laboratory Review Checkli	st: Reportable Data					
Lat	orato	ory Name: STL-Houston	RC Date: 06/26/06					
Pro	ject N	Name: 9003 N. MAIN	aboratory Job Number: 318054					
Rev	iewe	r Name: ERA	Prep Batch Number(s): 157626 (Water)-B	TEX			,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	
#1	A ²	Description		Yes	No	NA ³	NR ⁴	ER#5
S1	01	Initial calibration (ICAL)						77.0
	1	Were response factors and/or relative response factors for each	th analyte within OC limits?	X	1			
	1	Were percent RSDs or correlation coefficient criteria met?		X				1
		Was the number of standards recommended in the method us	ed for all analytes?	X				
		Were all points generated between the lowest and highest star		X				1
		Are ICAL data available for all instruments used?		X				
		Has the initial calibration curve been verified using an approp	oriate second source standard?	X				
S2	OI	Initial and continuing calibration verification (ICCV and						
		Was the CCV analyzed at the method-required frequency?		X				
		Were percent differences for each analyte within the method-	required QC limits?	X	1			
		Was the ICAL curve verified for each analyte?		X	T			
		Was the absolute value of the analyte concentration in the inc	organic CCB < MDL?		Ī	X		
S3	0	Mass spectral tuning:						
		Was the appropriate compound for the method used for tuning	g?			X		
		Were ion abundance data within the method-required QC lim			1	X		
S4	0	Internal standards (IS):		1.4			10 10 1	
		Were IS area counts and retention times within the method-re	quired QC limits?	X				
S5	OI	Raw data (NELAC section 1 appendix A glossary, and sec	tion 5.12 or ISO/IEC 17025 section		1 1 14		7 T Z	
		Were the raw data (for example, chromatograms, spectral data	a) reviewed by an analyst?	X				
		Were data associated with manual integrations flagged on the	raw data?	X				
S6	0	Dual column confirmation			1	4.4	1111	100
		Did dual column confirmation results meet the method-requir	ed QC?		X			4
S7	0	Tentatively identified compounds (TICs):						
		If TICs were requested, were the mass spectra and TIC data s	ubject to appropriate checks?			X		
S8	I	Interference Check Sample (ICS) results:		1 7 7	1 5,21 5,	177	N 10.	
		Were percent recoveries within method QC limits?				X		
S9	I	Serial dilutions, post digestion spikes, and method of stand	lard additions	1 1 1 1 1 1	19.35	44	1.0 4 1.1	
		Were percent differences, recoveries, and the linearity within	the QC limits specified in the method?			X		
S10	OI	Method detection limit (MDL) studies		151.5	10 Sec.	1.5	134.12	
		Was a MDL study performed for each reported analyte?		X				
		Is the MDL either adjusted or supported by the analysis of DC	CSs?	X				
S11	OI	Proficiency test reports:		45.5	170.3	7.00	1.7	
		Was the laboratory's performance acceptable on the applicable	e proficiency tests or evaluation studies?	X			l	
S12	Ol	Standards documentation				1.41	nin .	1.3
		Are all standards used in the analyses NIST-traceable or obtain	ned from other appropriate sources?	X				
S13	OI	Compound/analyte identification procedures						
		Are the procedures for compound/analyte identification documents	mented?	X				
S14	OI	Demonstration of analyst competency (DOC)		3.7				<u> </u>
		Was DOC conducted consistent with NELAC Chapter 5C or l		X	<u> </u>			
	<u> </u>	Is documentation of the analyst's competency up-to-date and		X				
S15	OI	Verification/validation documentation for methods (NELA						1.1 1
		Are all the methods used to generate the data documented, ver	rified, and validated, where applicable?	Х				
S16	OI	Laboratory standard operating procedures (SOPs):			1,14	77 1 14		117.5
		Are laboratory SOPs current and on file for each method perfe	ormed?	X				
	L			<u> </u>			L	L

Items identified by the letter "R" should be included in the laboratory data package submitted to the TCEQ in the TRRP-required report(s).

Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.

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Appe	ndix A (cont'd): Laboratory Review	Checklist: Exception Reports			
Laborat	tory Name: STL-Houston	LRC Date: 06/26/06			
Project Name: 9003 N. MAIN		Laboratory Job Number: 318054			
Reviewer Name: ERA		Prep Batch Number(s): 157626 (Water)-BTEX			
ER# ¹	DESCRIPTION				
1	The temperature of the cooler received b	y the laboratory on 06/21/06 was above the acceptable range of 2.0-6.0 °C.			
2		veries on both columns in sample 318054-4 were above acceptance limits			
3	The laboratory selected another client's s	ample to perform as the MS/MSD.			
4	The benzene RPD between the two columns in sample 318054-2 was >40%. Since anomalies were present, the lower of the two results was reported.				

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked on the LRC)

Lab	orato	ry Name; STL-Houston LRC	C Date: 06/26/06					
Proi	ect N	lame: 9003 N. MAIN Lab	oratory Job Number: 318054	••••••				
,				nton) E	·	aldah	do	
#1	A ²		Batch Number(s): 157516 (soil and W					ER#5
#	A ^s	Description12		Yes	No	NA.	NK	EK#
		Chain-of-custody (C-O-C)						
RI	OI	Did samples meet the laboratory's standard conditions of samp			<u>X</u>	ļ		1
		Were all departures from standard conditions described in an ex-	xception report?	X		<u> </u>		
R2	OI	Sample and quality control (QC) identification		1		L		
		Are all field sample ID numbers cross-referenced to the laborat		X		ļ		
	- T	Are all laboratory ID numbers cross-referenced to the correspo	nding QC data?	X		ļ		ļ
R3	OI					ļ	ļ	ļ
		Were all samples prepared and analyzed within holding times?		X				
		Other than those results < MQL, were all other raw values brace	keted by calibration standards?	X		 -		
		Were calculations checked by a peer or supervisor? Were all analyte identifications checked by a peer or superviso	w?	$\frac{\Lambda}{X}$				
		Were an analyte identifications checked by a peer or superviso. Were sample quantitation limits reported for all analytes not de		$\frac{X}{X}$				
		Were all results for soil and sediment samples reported on a dry		X				
		Were % moisture (or solids) reported for all soil and sediment s		$\frac{\hat{x}}{x}$				
		If required for the project, TICs reported?	samprest	+^+		Х		
R4	0	Surrogate recovery data		++		<u>^`</u>		
	<u> </u>	Were surrogates added prior to extraction?		$ \mathbf{x} $			-	·
		Were surrogate percent recoveries in all samples within the lab	oratory OC limits?	$\frac{1}{x}$				
R5	OI	Test reports/summary forms for blank samples	1 1					
		Were appropriate type(s) of blanks analyzed?		X				
		Were blanks analyzed at the appropriate frequency?		X				
		Were method blanks taken through the entire analytical process	s, including preparation and, if	X				
		applicable, cleanup procedures?						
		Were blank concentrations < MQL?		X				
R6	OI	Laboratory control samples (LCS):						
		Were all COCs included in the LCS?		X				
		Was each LCS taken through the entire analytical procedure, in	cluding prep and cleanup steps?	X				
		Were LCSs analyzed at the required frequency?		X				
		Were LCS (and LCSD, if applicable) %Rs within the laborator		X				***********
		Does the detectability data document the laboratory's capability	y to detect the COCs at the MDL used			X		
		to calculate the SQLs?		↓				
	<u> </u>	Was the LCSD RPD within QC limits?		11		X		
R7	OI	Matrix spike (MS) and matrix spike duplicate (MSD) data	1 Lang	 ,, 				
		Were the project/method specified analytes included in the MS	and MSD?	X				
		Were MS/MSD analyzed at the appropriate frequency? Were MS (and MSD, if applicable) %Rs within the laboratory of	OC 1::	X			V	
			QC limits?	+-+			XX	2
R8	OI	Were MS/MSD RPDs within laboratory QC limits? Analytical duplicate data		+-+			- 	
10	01	Were appropriate analytical duplicates analyzed for each matrix	······································	╂╼╂		х		
		Were analytical duplicates analyzed at the appropriate frequence		+-+		X		
		Were RPDs or relative standard deviations within the laboratory		╁╌┼		$\frac{\Lambda}{X}$		
R9	OI	Method quantitation limits (MQLs):	y QC imms:	\vdash		^		
		Are the MQLs for each method analyte included in the laborato	ry data package?	$ \mathbf{x} $				
		Do the MQLs correspond to the concentration of the lowest nor		$\frac{x}{x}$		-		
1		Are unadjusted MQLs included in the laboratory data package?		$\frac{1}{X}$	\neg	$\neg \uparrow$	$\neg \neg$	
R10		Other problems/anomalies		T		一	$\neg \dagger$	
		Are all known problems/anomalies/special conditions noted in t	this LRC and ER?	x			$\neg \uparrow$	
		Were all necessary corrective actions performed for the reported		X				
		Was applicable and available technology used to lower the SQI	to minimize the matrix interference	X	\neg			

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ΑI

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		ndix A (cont'd): Laboratory Review Checklist: R						
			te: 06/26/06					
Pro	ject N	Name: 9003 N. MAIN Laborat	ory Job Number: 318054					
Rev	iewe	er Name: JPS Prep Ba	tch Number(s): 157516 (soil and V	Wate	r)-Fo	rmald	ehyde	
#1	I A ²	Description		Yes		NA ³	NR ⁴	ER#
		Initial calibration (ICAL)	<u>,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,</u>			15 (15)	16,47	11.11
<u> </u>	 ``	Were response factors and/or relative response factors for each analy	te within OC limits?	X		 	 	
		Were percent RSDs or correlation coefficient criteria met?		X				
		Was the number of standards recommended in the method used for a	l analytes?	X	 		1	
		Were all points generated between the lowest and highest standard us		X				1
		Are ICAL data available for all instruments used?		X				
		Has the initial calibration curve been verified using an appropriate se	cond source standard?	X			<u> </u>	1
S2	OI	Initial and continuing calibration verification (ICCV and CCV) a			11111	11111	1.50	
	1	Was the CCV analyzed at the method-required frequency?		Х				1
		Were percent differences for each analyte within the method-required	OC limits?	Х				1
]	Was the ICAL curve verified for each analyte?		X	 	<u> </u>	<u> </u>	1
		Was the absolute value of the analyte concentration in the inorganic	CCB < MDL?		<u> </u>	X		
S3	0	Mass spectral tuning:					1	
	Ť	Was the appropriate compound for the method used for tuning?				Х	 	1
		Were ion abundance data within the method-required QC limits?			l	X	1	
S4	0	Internal standards (IS):		1 4- 13	3, 4,			100
	<u> </u>	Were IS area counts and retention times within the method-required	OC limits?		·	X		
S5	OI	Raw data (NELAC section 1 appendix A glossary, and section 5.1		15.1				
	<u>~ - </u>	Were the raw data (for example, chromatograms, spectral data) revie		X				
		Were data associated with manual integrations flagged on the raw da		X				
S6	0	Dual column confirmation						
	 	Did dual column confirmation results meet the method-required QC?				X		
S7	o	Tentatively identified compounds (TICs):						
	 	If TICs were requested, were the mass spectra and TIC data subject to	appropriate checks?			X	<u> </u>	
S8	1	Interference Check Sample (ICS) results:		7				1
	† 	Were percent recoveries within method QC limits?				X		
S9	I	Serial dilutions, post digestion spikes, and method of standard ad	ditions	Spira	354	2.54	1966	
	Ť-	Were percent differences, recoveries, and the linearity within the QC	limits specified in the method?			X		
S10	OI	Method detection limit (MDL) studies		100	100	200	1000	
	<u> </u>	Was a MDL study performed for each reported analyte?		X				1
	 	Is the MDL either adjusted or supported by the analysis of DCSs?		X				1
S11	OI	Proficiency test reports:		124.	75.0	diffracti	4445	VAX.
		Was the laboratory's performance acceptable on the applicable profic	iency tests or evaluation studies?	X				
S12	OI	Standards documentation		1.7				
	-	Are all standards used in the analyses NIST-traceable or obtained fro	m other appropriate sources?	X				1
S13	OI	Compound/analyte identification procedures						
	1	Are the procedures for compound/analyte identification documented?		X				
S14	OI	Demonstration of analyst competency (DOC)						l
	T	Was DOC conducted consistent with NELAC Chapter 5C or ISO/IEC	24?	X		+		
	1	Is documentation of the analyst's competency up-to-date and on file?		X				
S15	OI	Verification/validation documentation for methods (NELAC Char		11.	47,74		. 1. T. T.	1
	† **	Are all the methods used to generate the data documented, verified, a		X				T
\$16	OI	Laboratory standard operating procedures (SOPs):			. 15.	145.1	1 fe i	
	10,	Are laboratory SOPs current and on file for each method performed?		X			h	
	L	The modulatory box sources and on the for each medica performed					<u> </u>	<u></u>

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Apper	ndix A (cont'd): Laboratory Revie	ew Checklist: Exception Reports
Laborat	ory Name: STL-Houston	LRC Date: 06/26/06
Project	Name: 9003 N. MAIN	Laboratory Job Number: 318054
Review	er Name: JPS	Prep Batch Number(s): 157516 (soil and Water)-Formaldehyde
ER#1	DESCRIPTION	
1	The temperature of the cooler received	d by the laboratory on 06/21/06 was above the acceptable range of 2.0-6.0 °C.
2	The laboratory selected another client	's sample to perform as the MS/MSD.

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked on the LRC)

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CHAIN OF CUSTODY RECORD

Gustomet Information	Q.			; ;		6
1	PROJECT NAME	31-5.6.10.11	9003 N W .	76.0 76.0		,
0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0		IE	'V', 'V'\a'W	O C: 8315 F01	B.TPH TX 1005 C:8315 Formaldehyde	
COMPANY ATC Associates, inc.	BILL TO	ATC Associates, Inc.	,		,	
SEND REPORT TO Patrick Duoraczyk	INVOICE ATTN	Patrick Dworaczyk			solis bry meighter	P karina kalauna kar
AODRESS 3928 Blaebonnet Drive	ADDRESS 39	3928 Bluedonnet Orive		т _ ·		
				- × -		
				J Z Z	0,8054	
GITWSTATEZIP Stafford, Il 77477	CITY/STATEZIP	Stafford, TX 77477		2 O a	7 0 1	************
PHONE: 281-240-0154	PHONE 28	281-240-0154		201		
FAX 281-240-8909	FAX. 8.28.	281-240-8909		rω		***************************************
SAMP NO SAMPLE DESCRIPTION	PRESERVE	F SAMPLE MATRIX SA	SAMPLE DATE SA	SAMPLE TIME # CONTAINER	ER A B C D E F G H I J K L M NO	POR
3-1 12-14		7.98	90-12-9	950		
B-2 20-22			╂	1100		
B3 20-22		Sp;		11.55		
<u>B.1</u>		Water		arci	7,7,7	
[የጋ		Water	<u> </u>	1245	N.V.	
3-3		Water		001		
Sampler:	Shipment Method: Ares of	Airbill	No.:		Required TurnAround: 5	
1. Relinquished By Reference 1	Date 2. Relinquished By:		Date	3. Relinquished By:		Date
Company Name:	Time Company Name:		Time	Company Name:	<u></u>	Time
1. Received By: GMTD	pate 2. Received By:		Date	3. Received By:	0	Date
Company Name:	Time, 24 Company Name:		Time	Company Name:		Time
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713-690-4444 RAX 713-690-5646 Bouston, TX 77040 Severn Trent Laboratories 6310 Rothway Brive

rpjsckl	Job Sample Receipt Checklist Report	V2
Customer Job ID:	n.: 57216 Check List Number.: 1 Description.: Job Check List Date.: 06/21/2006 ject Description.: TRRP Project , Inc. Contact.: Patrick Dwora	Date of the Report: 06/21/2006 Project Manager: daj1 czyk
Questions ?	(Y/N) Comments	
Chain of Custody Received?	Y	
If "yes", completed properly	? Y	
Custody seal on shipping contain	ner? N	
If "yes", custody seal intac	t?	
Custody seals on sample contain	ers? N	
If "yes", custody seal intac	t?	
Samples chilled?	N see src	
Temperature of cooler acceptabl	e? (4 deg C +/- 2). N 18.6	
If "no", is sample an air ma	trix?(no temp req.) N	
Thermometer ID	Y 437	2 /
Samples received intact (good c	ondition)? Y	()(
Volatile samples acceptable? (n	o headspace) Y	10-21-6
Correct containers used?	Y	10-1/1
Adequate sample volume provided	17Y	V
Samples preserved correctly?	,,,,,,Y	
Samples received within holding	y-time? Y y	
Agreement between COC and sampl	e labels? Y	
Radioactivity at or below backg	round levels? Y	
Additional		
Comments		
Sample Custodian Signature/Date	Y jac	

Page 1

V	4 Tre			- SAMPLE		CHECKLIS	r/, ,
CLIENT NAME:	/	<u> </u>		CARRIE	ER/DRIVER	NAME:	Client
PROJECT:	·	·	UN	PACKED BY:	Profession		
DATE RECEIVED:			***************************************		UNPACKE	ED STAMP:	
TOTAL # COOLER	S RECEIVED):					
1				OLER CHE	CKLIST		
COOLER ID	COC PRESENT	CUSTOD	Y TAPE	COOLER TEMP	THERM ID	TEMP BLK PRESENT	List Sample Bottles in Each Cooler if out of Temperature
	(Y/N)	PRESENT (Y/N)	INTACT (Y/N)	(°C)	טו	(Y/N)	out of remperature
Gray lihit,	y	C B/V	N	18.6	437	N	Chillippograss
	,	С	· · · · · · · · · · · · · · · · · · ·		·		11/5
		В					
		С					
C=COOLER B	= BOTTLES	В					
** SPECIFIC PROJEC VOLATILE HEADSF (If ANY headspace is p pH OF WATER SAM PRESERV H2SO4 (<2) HNO3 (<2) HCL (<2) (Not VOA NaOH – Cyanide (> NaOH/Zn Acetate – Other	PACE ACCEF resent, list det MPLES /ATION Vials)	PTABLE? Yes ails in INCONSI	No	NA	Je M N	umber of VO	**************************************
# OF NEAT BOTTLE	ES:				#	OF SOIL JAF	RS:
		INCONSIST	ENCIES (Place in Job	Notes as v	vell (CTRL F	-12)
PERSON CONTACT RESOLUTION	ED:			ACTION TAP		DATE:	
NOTES							
Project Manager					****	(Use b	ack of sheet if necessary)



7/10/2006 Date:

> B-1 12-14 Customer Sample ID:

Job Number: 318054

Date/Time Sampled 6/21/2006

09:50 13:31 Date/Time Received: 6/21/2006

Laboratory Sample ID: 318054-001

Sample Matrix Soil

			Value	S WIDT	MOIN	108		Analysis Date June	Batch	10	Vralyst
Method: SM-2540 G Mod., Soil											
% Solids	NA	82.2				aller en Production en	%	6/22/2006 16:30	157440	1.00	ydh
Moisture	MOIST	17.8				and the second second	%	6/22/2006 16:30	157440	1.00	yps
Method: SW-846 8021B, Soil											
Benzene	71-43-2	5.60	D D	4.60	10.0	5.60	ug/Kg	6/23/2006 20:50	157581	1.00	era
Ethylbenzene	100-41-4	7.42	Þ	6.10	10.0	7.42	ug/Kg	6/23/2006 20:50	157581	1.00	era
m,p-Xylene	MPXYLENE	15.4	Þ	12.7	20.0	15.4	ug/Kg	6/23/2006 20:50	157581	1.00	era
o-Xylene	95-47-6	7.86	Ð	6.46	10.0	7.86	ug/Kg	6/23/2006 20:50	157581	1.00	ега
Toluene	108-88-3	7.74	D	6.36	10.0	7.74	ug/Kg	6/23/2006 20:50	157581	1.00	era
Xylenes (total)	1330-20-7	23.3	D	19.1	30.0	23.3	ug/Kg	6/23/2006 20:50	157581	1.00	era
Method: SW-846 8315, Soil											
Formaldehyde	0-00-09	2520		2.87	100	6.69	ug/Kg	6/26/2006 11:56	157630	1.00	jps
Method: SW846 8315, Water											
Solid Phase Extraction	NA	Complete					N/A	6/23/2006 08:00	157516	1.00	oue
									•		***************************************

Form I



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-1 12-14

Date/Time Sampled: 6/21/2006 09:50

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-001

Sample Matrix Soil

D. E. H. Analysti		1.00 mep	I.00 mep	1.00 mep	1.00 mep		
Batch		157465	157465	157465	157465		157296
AnalysisDareTime		6/22/2006 20:38	6/22/2006 20:38	6/22/2006 20:38	6/22/2006 20:38		6/21/2006 16:00
SUND			mg/Kg	mg/Kg	mg/Kg		N/A
NOS		13.8	13.8	6.57	13.8		
MOI		50.0	50.0	20.0	20.0		
MDI		11.4	11.4	5.40	11.4		
		þ	n	D	n		
NESC IN		13.8	13.8	6.57	13.8		Complete
		NA	NA	NA	NA		N A
	TNRCC 1005, Soil	Petroleum Hydrocarbons C12 - C28	Petroleum Hydrocarbons C28 - C35	Petroleum Hydrocarbons C6 - C12		Method: TNRCC TX-1005, Soil	TNRCC 1005 Extraction

Form I



Date: 7/10/2006

Customer Sample ID: B-2 20-22

Job Number: 318054

Date/Time Sampled 6/21/2006 11:00

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-002

Sample Matrix Soil

	Svo	NESS COL		MDE		108		Analysis Date mile	Batch	DE	Analyst
Method: SM-2540 G Mod., Soil											
% Solids	NA	85.2			-	:	%	6/22/2006 16:30	157440	1.00	sdħ
Moisture	MOIST	14.8					%	6/22/2006 16:30	157440	1.00	sdħ
Method: SW-846 8021B, Soil											
Benzene	71-43-2	72.0		4.60	10.0	5.40	ug/Kg	6/23/2006 21:30	157581	1.00	era
Ethylbenzene	100-41-4	785		6.10	10.0	7.16	ug/Kg	6/23/2006 21:30	157581	1.00	era
m,p-Xylene	MPXYLENE	1450		12.7	20.0	14,9	ug/Kg	6/23/2006 21:30	157581	1.00	era
o-Xylene	95-47-6	512		6.46	10.0	7.58	ug/Kg	6/23/2006 21:30	157581	1.00	era
Toluene	108-88-3	155		6.36	10.0	7.46	ug/Kg	6/23/2006 21:30	157581	1.00	era
Xylenes (total)	1330-20-7	1970		19.1	30.0	22.5	ug/Kg	6/23/2006 21:30	157581	1.00	era
Method: SW-846 8315, Soil											
Formaldehyde	20-00-0	575		2.87	100	67.3	ug/Kg	6/26/2006 12:12	157630	1.00	jps
Method: SW846 8315, Water											
Solid Phase Extraction	NA	Complete					N/A	6/23/2006 08:00	157516	1.00	enc

Form I



Date: 7/10/2006

Customer Sample ID: B-2 20-22

Job Number: 318054

Date/Time Sampled 6/21/2006 11:00

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-002

Sample Matrix Soil

COPER S						 		
Analyst.		mep	mep	mep	mep		lvp	
		1.00	1.00	1.00	1.00		1.00	
Batch		157465	157465	157465	157465		157296	
Dole Inc		6/22/2006 21:44	6/22/2006 21:44	6/22/2006 21:44	6/22/2006 21:44		6/21/2006 16:00	
Affilia		6/22/	6/22/	6/22/	6/22/		6/21/	
A VIOLET SOLUTION DE LA MINE AMBESTS DATE MANAGEMENT		mg/Kg	mg/Kg	mg/Kg	mg/Kg		N/A	
SOI		13.3	13.3	6.32	13.3			
NO		20.0	50.0	50.0	50.0			
MDI		11.4	11.4	5.40	11.4			
Ø.		<u> </u>	<u> </u>	<u> </u>	Þ			
RESULT OF		13.3	13.3	6.32	13.3		Complete	
English Substitute		NA	NA	NA	NA		NA V	
	Method: TNRCC 1005, Soil	Petroleum Hydrocarbons C12 - C28	Petroleum Hydrocarbons C28 - C35	Petroleum Hydrocarbons C6 - C12	Petroleum Hydrocarbons C6 - C35	Method: TNRCC TX-1005, Soil	TNRCC 1005 Extraction	
	Meth	Petrole	Petrole	Petrol	Petrole	Meth	TINEC	

Form I



Date: 7/10/2006

USECMEN PARCEASSOCIAL E

Job Number: 318054

Customer Sample ID: B-3 20-22

Date/Time Sampled: 6/21/2006 11:55

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-003

Sample Matrix Soil

	To AS T	RESULT		JUN	MOLE	SOL		Makis Bate mine	Batch		Analysi
Method: SM-2540 G Mod., Soil											
% Solids	NA	85.3					%	6/22/2006 16:30	157440	1.00	sdh
Moisture	MOIST	14.7		to reconstruction			%	6/22/2006 16:30	157440	1.00	ydh
Method: SW-846 8021B, Soil											
Benzene	71-43-2	5.39	5	4.60	10.0	5.39	ug/Kg	6/23/2006 22:30	157581	1.00	era
Ethylbenzene	100-41-4	7.15	D D	6.10	10.0	7.15	ug/Kg	6/23/2006 22:30	157581	1.00	era
m,p-Xylene	MPXYLENE	14.9		12.7	20.0	14.9	ug/Kg	6/23/2006 22:30	157581	1.00	era
o-Xylene	95-47-6	7.57		6.46	10.0	7.57	ug/Kg	6/23/2006 22:30	157581	1.00	era
Toluene	108-88-3	7.45	D	6.36	10.0	7.45	ug/Kg	6/23/2006 22:30	157581	1.00	era
Xylenes (total)	1330-20-7	22.4	n	19.1	30.0	22.4	ug/Kg	6/23/2006 22:30	157581	1.00	era
Method: SW-846 8315, Soil											
Formaldehyde	20-00-0	357		2.87	100	67.3	ug/Kg	6/26/2006 12:27	157630	1.00	sdí
Method: SW846 8315, Water											
Solid Phase Extraction	NA	Complete					N/A	6/23/2006 08:00	157516	1.00	enc

Form I



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-3 20-22

Date/Time Sampled: 6/21/2006

11:55

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-003

Sample Matrix Soil

Set State		0 mep	0 mep	0 mep	0 mep		0 Ivp	
G		1.00	1.00	1.00	1.00		1.00	
Batch		157465	157465	157465	157465		157296	
Analysis Date Illines Barel		6/22/2006 22:17	6/22/2006 22:17	6/22/2006 22:17	6/22/2006 22:17		6/21/2006 16:00	
Analysis		6/22/20	6/22/20	6/22/20	6/22/20		6/21/20	
		mg/Kg	mg/Kg	mg/Kg	mg/Kg		N/A	
TOS		13.3	13.3	6.33	13.3			
TON!		50.0	50.0	50.0	50.0			
VOI		11.4	11.4	5.40	11.4			
5)								·
0		Ω	Þ		ם		<u></u>	
KESU		13.3	13.3	6.33	13.3		Complete	
		¥	4	∢	₹		¥	
		NA	NA	N A	NA		NA	
		C28	C35	212	335	Soil		
TO STOWER THOUSA	Method: TNRCC 1005, Soil	Petroleum Hydrocarbons C12 - C28	Petroleum Hydrocarbons C28 - C35	Petroleum Hydrocarbons C6 - C12	Petroleum Hydrocarbons C6 - C35	Method: TNRCC TX-1005, Soil	ď	
	3C 186	carbon	carbon	carbon	carbon:	XI X	TNRCC 1005 Extraction	
	TNRC	Hydroc	Hydro	Hydroc	Hydroc	TNRC	05 Exi	
	iod:	leum l	leum l	leum]	leum J	:poq:	CC 10	
	Meti	Petro	Petro	Petro	Petro	Met	TAR	

Form I



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-1

Date/Time Sampled: 6/21/2006 12:10

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-004

Sample Matrix Water

THE STANFORD STANFORD		RESULT	9 FE A.G		Moj	Sol		Analysis Dale Time	Barch	. TO	Malya
Method: SW-846 8021B, Water											
Benzene	71-43-2	25.6	·	0.170	0.500	0.170	ng/L	6/26/2006 10:20	157626	1.00	era
Ethylbenzene	100-41-4	131		0.230	0.500	0.230	T/gn	6/26/2006 10:20	157626	1.00	era
m,p-Xylene	MPXYLENE	297		0.370	1.00	0.370	ng/L	6/26/2006 10:20	157626	1.00	era
o-Xylene	95-47-6	6.66		0.190	0.500	0.190	T/Sn	6/26/2006 10:20	157626	1.00	ета
Toluene	108-88-3	7.05		0.190	0.500	0.190	ng/L	6/26/2006 10:20	157626	1.00	ета
Method: SW846 8315, Water											
Solid Phase Extraction	NA	Complete					N/A	6/23/2006 08:00	157516	1.00	enc
Method: SW-846 8315, Water											
Formaldehyde	20-00-0	27.6		2.87	5.00	3.38	T/fin	6/26/2006 12:58	157630	1.00	sdí
Method: TNRCC 1005, Water											
Petroleum Hydrocarbons C12 - C28	NA	0.830		0.870	2.00	0.830	mg/L	6/23/2006 23:03	157591	1.00	mep
Petroleum Hydrocarbons C28 - C35	NA	0.830		0.870	5.00	0.830	mg/L	6/23/2006 23:03	157591	1.00	шер
Petroleum Hydrocarbons C6 - C12	NA	0.480	ם	0.500	2.00	0.480	mg/L	6/23/2006 23:03	157591	1.00	mep
Petroleum Hydrocarbons C6 - C35	NA	0.830	D	0.870	5.00	0.830	mg/L	6/23/2006 23:03	157591	1.00	dəm
-											

Form I

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6310 Rothway Drive • Houston, TX 77040 • Tel: 713 690 4444 • Fax: 713 690 5646 • www.stl-inc.com



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-1

Date/Time Sampled 6/21/2006 12:10

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-004

Sample Matrix Water

Method: TNRCCTX-1005, Water	TNRCC 1005 Extraction
	ž
	Complete
	A/N
	6/22/2006 14:30
	157449
	1.00 lvp

Form I



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-2.

Date/Time Sampled: 6/21/2006 12:45

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-005

Sample Matrix Water

THE WILLIAM DESCRIPTION OF THE PROPERTY OF THE		THE SERVICE OF		KCIIV. IIIO	IOM	NOS.	SHAULE	Analysis Date Trine	Batch	DIC	A CHELLY ST
Method: SW-846 8021B, Water											
Benzene	71-43-2	0.170	n	0.170	0.500	0.170	ng/L	6/26/2006 10:40	157626	1.00	era
Ethylbenzene	100-41-4	0.670		0.230	0.500	0.230	ng/L	6/26/2006 10:40	157626	1.00	era
m,p-Xylene	MPXYLENE	1.81		0.370	1.00	0.370	ng/L	6/26/2006 10:40	157626	1.00	era
o-Xylene	95-47-6	0.560	***************************************	0.190	0.500	0.190	J/gn	6/26/2006 10:40	157626	1.00	era
Toluene	108-88-3	0.190	n	0.190	0.500	0.190	ug/L	6/26/2006 10:40	157626	1.00	era
Method: SW846 8315, Water											
Solid Phase Extraction	NA	Complete					N/A	6/23/2006 08:00	157516	1.00	enc
Method: SW-846 8315, Water											
Formaldehyde	20-00-0	46.4		2.87	5.00	3.38	ng/L	6/26/2006 13:13	157630	1.00	ips
Method: TNRCC 1005, Water											
Petroleum Hydrocarbons C12 - C28	NA	0.830	n	0.870	5.00	0.830	mg/L	6/23/2006 23:36	157591	1.00	mep
Petroleum Hydrocarbons C28 - C35	NA	0.830	n	0.870	2.00	0.830	mg/L	6/23/2006 23:36	157591	1.00	mep
Petroleum Hydrocarbons C6 - C12	NA	0.480	D	0.500	5.00	0.480	mg/L	6/23/2006 23:36	157591	1.00	mep
Petroleum Hydrocarbons C6 - C35	NA	0.830	ב	0.870	5.00	0.830	mg/L	6/23/2006 23:36	157591	1.00	mep
		****		•	-	-					_

Form I



7/10/2006 Date:

Job Number: 318054

Customer Sample ID:

12:45 Date/Time Sampled 6/21/2006

13:31 Date/Time Received 6/21/2006

Laboratory Sample ID: 318054-005

Sample Matrix Water

Contraction de la contraction		
		qv
300		1.00
Bates.		157449
THIC		
Spirit and		6/22/2006 14:30
		6/22
		N/A
		-
108		
Military		
)		
		Сотрієє
		СО
188		
		₹ Z
	Vater	
0.01	-1005, V	
	CCTX	TNRCC 1005 Extraction
	t: TNR	1005 E
	Methoc	TNRCC
	1 T T	

Form I



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-3

Date/Time Sampled 6/21/2006 13:00

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-006

Sample Matrix Water

HAND TOURS IN SERVICE		KASIII T		GIV NO	MOI	10S	TINITS F	AnalysisDaterume		110	finalyst
Method: SW-846 8021B, Water											
Benzene	71-43-2	0.170	<u></u>	0.170	0.500	0.170	ng/L	6/26/2006 11:57	157626	1.00	era
Ethylbenzene	100-41-4	0.230	D	0.230	0.500	0.230	ng/L	6/26/2006 11:57	157626	1.00	era
m,p-Xylene	MPXYLENE	0.370	D	0.370	1.00	0.370	ng/L	6/26/2006 11:57	157626	1.00	ета
o-Xylene	95-47-6	0.190	_	0.190	0.500	0.190	T/Sn	6/26/2006 11:57	157626	1.00	cra
Toluene	108-88-3	0.190	D	0.190	0.500	0.190	ug/L	6/26/2006 11:57	157626	1.00	era
Method: SW846 8315, Water											
Solid Phase Extraction	NA	Complete		tan an kast			N/A	6/23/2006 08:00	157516	1.00	enc
Method: SW-846 8315, Water											
Formaldehyde	20-00-0	9.80		2.87	5.00	3.54	ng/L	6/26/2006 13:29	157630	1.00	jps
Method: TNRCC 1005, Water											
Petroleum Hydrocarbons C12 - C28	NA	0.840	n n	0.870	2.00	0.840	mg/L	6/24/2006 00:09	157591	1.00	шер
Petroleum Hydrocarbons C28 - C35	NA	0.840	Þ	0.870	5.00	0.840	mg/L	6/24/2006 00:09	157591	1.00	mcp
Petroleum Hydrocarbons C6 - C12	NA	0.480	n	0.500	5.00	0.480	mg/L	6/24/2006 00:09	157591	1.00	mep
Petroleum Hydrocarbons C6 - C35	NA	0.840)	0.870	5.00	0.840	mg/L	6/24/2006 00:09	157591	1.00	mep

Form I



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-3

Date/Time Sampled 6/21/2006 13:00

.Date/Time Received 6/21/2006 13:31

Laboratory Sample ID: 318054-006

Sample Matrix Water

lvp

1.00

157449

6/22/2006 14:30

N/A

Complete NA Method: TNRCC TX-1005, Water TNRCC 1005 Extraction

Form 1



QUALITY CONTROL RESULTS

Job Number.: 318054

Report Date.: 07/10/2006

CUSTOMER: ATC Associates, Inc.

PROJECT: 9003 N. MAIN

ATTN: Patrick Dworaczyk

Mε	est Method thod Descrip arameter	otion,: Moi	sture (Total +	Fixed Solids,					st: sdh Code.: %SOLID	
QC	Lab ID	Reagent	QC Result	QC Result	True Value	Orig. Value	Calc. Result *	Limits	F Date	Tim
DU	318054-3		85.8368	<u> </u>		85.3351	0.6	10.0	06/22/2006	163
UC	318174-6		83.8722			83.3475	0.6	10.0	06/22/2006	163
U	318107-4		98.3661			98.3314	0.0	10.0	06/22/2006	163
DU	318174-16		85.3161			85.0272	0.3	10.0	06/22/2006	163
MB	15744021		0.0000						06/22/2006	163
MB	15744021		0.0000						06/22/2006	163
MB	15744021		0.0000						06/22/2006	
DU	318100-8		79.4863			78.5088	1.2	10.0	06/22/2006	163
DU	318146- 1		15.6110			15.7773	1.1	10.0	06/22/2006	163
Me Pe	irameter	otion.: Moi	sture (Total + sture	Fixed Solids,	Batch(s): 157440		Test	st: sdh Code.: MOIST	7:-
SC	Lab ID	Reagent	QC Result	QC Result	True Value	Orig. Value	Calc. Result *	Limits	F Date	Tin
U	318174-16		14.6839			14.9728	1.9	10.0	06/22/2006	
U	318174-6		16.1278			16.6525	3.2	10.0	06/22/2006	
UC	318100-8		20.5137			21.4912	4.7	10.0	06/22/2006	
U	318107-4		1.6339			1.6686	2.1	10.0	06/22/2006	
UC	318054-3		14.1632			14.6649	3.5	10.0	06/22/2006	
บเ	318146-1		84.3890			84.2227	0.2	10.0	06/22/2006	4/"



m,p-Xylene, Soil
o-Xylene, Soil
Xylenes (total), Soil
Total BTEX, Soil
Tert-Butyl Methyl Ether Column B, Soil

Methyl tert-Butyl ether, Soil

Benzene, Soil Toluene, Soil Ethylbenzene, Soil

Benzene Column B, Soil

Job Number.: 318054	QUALITY	CONTRO	. RESULI		eport Date.: 07	7/10/2006	
CUSTOMER: ATC Associates, Inc.	PROJE	CT: 9003 N. MA	i N	A	TTN: Patrick Dv	ioraczyk	
QC Type Description		Reag. Cod	e Lab	ID	Dilution Factor	Date	Time
Test Method: SW-846 80218 Method Description.: GC Volatile Organi	cs		: uç): 157581 1		Anal	yst: era	
LCS Laboratory Control Sample		BX\$0612060	157581-	ı		06/23/200	1709
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Va	lue Calc. Res	sult * Limi	ts F
Methyl tert-Butyl ether, Soil Benzene, Soil Toluene, Soil Ethylbenzene, Soil m,p-Xylene, Soil o-Xylene, Soil Xylenes (total), Soil Total BTEX, Soil Tert-Butyl Methyl Ether Column B, Soil Benzene Column B, Soil Toluene Column B, Soil Ethylbenzene Column B, Soil ethylbenzene Column B, Soil o-Xylene Column B, Soil	46.6816 52.0285 52.1341 52.0014 109.893 55.4779 166.0887 322.2527 45.1083 49.6298 51.3092 51.3521 104.727 56.1957		50.00000 50.00000 50.00000 100.00000 150.00000 150.00000 50.00000 50.00000 50.00000 50.00000 50.00000		93.4 104.1 104.3 109.9 111.0 110.7 107.4 90.2 99.3 102.6 102.7 104.7	69- 70- 71- 70- 70- 70- 61- 69- 71- 71-	125 133 134 139 136 131 130 130 125 133 134 139 136
MB Method Blank			157581-	1		06/23/200	6 1730
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Va	ilue Calc. Re	sult * Limi	its F
Methyl tert-Butyl ether, Soil Benzene, Soil Toluene, Soil Ethylbenzene, Soil m,p-Xylene, Soil o-Xylene, Soil Yotal BTEX, Soil Tert-Butyl Methyl Ether Column B, Soil Benzene Column B, Soil Toluene Column B, Soil Ethylbenzene Column B, Soil m,p-Xylene Column B, Soil o-Xylene Column B, Soil	ND ND ND ND ND 0.0000 ND ND ND ND ND ND						
SB Spiked Blank		BXS061206C	157581+	1		06/23/200	06 1930
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Va	olue Calc. Re	sult * Lim	its F

Page 35	*	%=% REC.	R=RPD, A=AB	S Diff.,	D=% Diff.

ND

ND

ND

ND

ND

ND

ND

ND

0.0000

0.0000

93

101

102

100

105

106

110

107

91

94

50.000000

50.000000

50.000000

50.000000

100.000000

50.000000

150.000000 300.000000

50.000000

50.000000

30.0-130.0

30.0-130.0 30.0-130.0 30.0-130.0

30.0-130.0

30.0-130.0

30.0-130.0

30.0-130.0

30.0-130.0

30.0-130.0

46.6336

50.6572 51.1370

50.0466

53.1742

165.5110

320.0681

45.5755

47.0232

104.969



Job Number.: 318054	QUALITY	CONTROL	RESULT		Date.: 07/10)/2006	
CUSTOMER: ATC Associates, Inc.	PROJEC	T: 9003 N. MAIN		ATTN:			
QC Type Description		Reag. Code	Lab	ID Dilut	ion Factor	Date Time	<u>.</u>
SB Spiked:Blank		BXS061206C	157581-1			06/23/2006 193	50
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Toluene Column B, Soil Ethylbenzene Column B, Soil m,p-Xylene Column B, Soil o-Xylene Column B, Soil	48.2890 52.7629 100.260 60.5420		50.000000 50.000000 100.000000 50.000000	ND ND ND ND	97 106 100 121	30.0-130.0 30.0-130.0 30.0-130.0 30.0-130.0	
SBD Spiked Blank Duplicate		BX\$061206C	157581-1			06/23/2006 195	. 0
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	: * Limits	F
Methyl tert-Butyl ether, Soil	49.6281	46.6336	50.000000	ND	99.3 6.2	30-130 20	
Benzene, Soil	54.0149	50.6572	50.000000	ND	108.0 6.4	30-130 20	
Toluene, Soil	54.0783	51.1370	50.000000	ND	108.2 5.6	30-130 20	
Ethylbenzene, Soil	54.2439	50.0466	50.000000	ND	108.5 8.0	30-130 20	
m,p-Xylene, Soil	113.256	104.969	100.000000	ND	113.3	30-130 20	
o-Xylene, Soil	56.5970	53.1742	50.000000	ND	7.6 113.2	30-130	
Xylenes (total), Soil	169.8530	165.5110	150.000000	0.0000	6.2 113.2	20 30-130	
Total BTEX, Soil	332.1901	320.0681	300.000000	0.0000	2.6 110.7	20 30-130	
Tert-Butyl Methyl Ether Column B, Soil	48.6404	45.5755	50.000000	ND	3.7 97.3	20 30-130	
Benzene Column B, Soil	50.4222	47.0232	50.000000	ND	6.5 100.8	20 30-130	
Toluene Column B, Soil	51.9796	48.2890	50.000000	ND	7.0 104.0	20 30-130	
Ethylbenzene Column B, Soil	53.2134	52.7629	50.000000	ND	7.4 106.4	20 30-130	
m,p-Xylene Column B, Soil	107.819	100.260	100.000000	ND	0.9 107.8	20 30-130	
o-Xylene Column B, Soil	55.7559	60.5420	50.000000	ND	7.3 111.5	20 30-130	
					8.2	20	
LCS Leberatory Control Sample		BXS0621068	157626-1			06/26/2006 072	29
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Methyl tert-Butyl ether, Water Benzene, Water Toluene, Water Ethylbenzene, Water m,p-Xylene, Water o-Xylene, Water Xylenes (total), Water Total BIEX, Water	49.1416 48.6675 49.1170 48.4982 99.1402 49.7578 148.8980 295.1807		50.000000 50.000000 50.000000 50.000000 100.000000 50.000000 150.000000 50.000000		98.3 97.3 98.2 97.0 99.1 99.5 99.3 98.4 97.7	76-123 72-134 76-131 75-131 75-130 74-129 70-130 76-123	
Tert-Butyl Methyl Ether Column B, Water Benzene Column B, Water	48.8600 47.5087		50.000000		95.0	76-123 72-134	

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* %=% REC, R=RPD, A=ABS Diff., D=% Diff.



	Job Number.: 318054	QUALITY	CONTROL	RESULT		Date.: 07/10	0/2006
CUSTOMER: AT	C Associates, Inc.	PROJEC	T: 9003 N. MAI	.	ATTN:		
QC Type	Description		Reag. Code	Lab I	ID Dilut	ion Factor	Date Time
LCS	Laboratory Control Sample		BX\$062106B	157626-1			06/26/2006 0729
Param	eter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	t * Limits F
Toluene Column Ethylbenzene C m,p-Xylene Col o-Xylene Colum	olumn B, Water umn B, Water	48.5180 47.7200 98.3282 48.3311		50.000000 50.000000 100.000000 50.000000		97.0 95.4 98.3 96.7	76-131 75-131 75-130 74-129
MB	Method Blank			157626-1			06/26/2006 0749
Param	eter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	t * Limits F
Benzene, Water Toluene, Water Ethylbenzene, m,p-Xylene, Water Xylenes (total Total BTEX, Water Tert-Butyl Met Benzene Column Toluene Column	Water iter ir), Water iter hyl Ether Column B, Water i B, Water i B, Water olumn B, Water umn B, Water	ND ND ND ND O.0000 O.0000 ND ND ND ND ND ND ND ND ND ND ND ND ND					
МВ	Method Blank			157626-1	20.0	00	06/26/2006 0819
Param	meter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Resul	t * Limits F
Benzene, SPLP Toluene, SPLP Ethylbenzene, m,p-Xylene, SF o-Xylene, SPLF Xylenes (total Total BTEX, SF	PLP), SPLP thyl Ether Column B, SPLP n B, SPLP n B, SPLP column B, SPLP .umn B, SPLP	ND ND ND ND ND ND O.0000 O.0000 ND ND ND ND ND ND ND ND ND ND					
MS	Matrix Spike		BX\$062306B	317399-1	20.0	00	D6/26/2006 D940
Param	neter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Resul	
Methyl tert-Bu Benzene, SPLP Toluene, SPLP Ethylbenzene,	utyl ether, SPLP	50.4870 43.9578 43.9321 43.0510		50.000000 50.000000 50.000000 50.000000	ND ND ND	88 88 86	70-130 70-130 70-130 70-130
			Page 37	* %=% REC,	R=RPD, A=ABS	Diff., D=% D	iff.



	Job Number.: 318054	QUALITY	CONTROL	RESULT		Date.: 07/10	/2006	
CUSTOMER: AT	C Associates, Inc.	PROJE	CT: 9003 N. MAI	N	ATTN:			
QC Type	Description		Reag. Code	Lab	ID Dilut	ion Factor	Date Time	e
MS	Matrix Spike		BXS062306B	317399-1	20,00	0	06/26/2006 094	40
Param	neter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
m,p-Xylene, SP o-Xylene, SPLP Xylenes (total Total BTEX, SP Tert-Butyl Met Benzene Column Toluene Column Ethylbenzene C m,p-Xylene Colum), SPLP LP thyl Ether Column B, SPLP B B, SPLP B B, SPLP Column B, SPLP cumn B, SPLP	89.7171 44.1903 133.9074 264.8483 50.6467 42.8537 42.8234 42.4284 87.2334 42.8149		100.000000 50.000000 150.000000 50.000000 50.000000 50.000000 50.000000 100.000000	ND 0.0000 0.0000 1.78440 ND ND ND	90 88 89 88 98 86 86 85 87	70-130 70-130 70-130 70-130 70-130 70-130 70-130 70-130 70-130	
MSD	Matrix Spike Duplicate		BXS062306B	317399-1	20,00	0	06/26/2006 100	00
Param	neter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Methÿl tert-Bu	ityl ether, SPLP	52.0804	50.4870	50.000000	1.37053	101.4 3.2	70-130 20	
Benzene, SPLP		44.1746	43.9578	50.000000	ND	88.3 0.5	70-130 20	
Toluene, SPLP		43.9849	43.9321	50.000000	ND	88.0	70-130	
Ethylbenzene,	SPLP	43.3784	43.0510	50.000000	ND	0.1 86.8	20 70-130	
m,p-Xylene, SP	PLP	90.1354	89.7171	100.000000	ND	0.8 90.1	20 70-130	
o-Xylene, SPLP	•	44.5674	44.1903	50.000000	ND	0.5 89.1	20 70-130	
Xylenes (total	.), SPLP	134.7028	133.9074	150.000000	0.0000	0.8 89.8	20 70-130	
Total BTEX, SP	•	266.2407	264.8483	300.000000	0.0000	0.6 88.7	20 70-130	
•	hyl Ether Column B, SPLP	52.1440	50.6467	50.000000	1.78440		20 70-130	
Benzene Column		42.8282	42.8537	50.000000	ND	3.0 85.7	20 70-130	
Toluene Column	•	42.7408	42.8234	50.000000	ND	0.1 85.5	20 70-130	
Ethylbenzene C	•	42.3851	42.4284	50.000000	ND	0.2 84.8	20 70-130	
m,p-Xylene Col	·	87,2536	87.2334	100.000000		0.1 87.3	20 70-130	
o-Xylene Colum	-	42.9718	42.8149	50.000000		0.0 85.9 0.4	20 70-130 20	



							•	
	Job Number.: 318054	QUALITY	CONTROL	. RESULT		t Date.: 07/10/	/2006	
CUSTOMER: A	TC Associates, Inc.	PROJEC	CT: 9003 N. MAI	N	ATTN:			
QC Type	Description		Reag. Code	e Lab	ID Dilu	tion Factor	Date Ti	me
	: TNRCC 1005 ription.: Direct Analytical	TPH Method TX 100		: mg/		Analyst	: тер	
LCD	Laboratory Control Sample	Dupticate	GCL051906	157296-1			06/21/2006 2	012
Parar	meter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hydi	rocarbons C6 - C12, Soil	236.881	234.799	250.000000	ND	95	70-130	
Petroleum Hyd	rocarbons C12 - C28, Soil	275.306	281.051	250.000000	ND	0.9 110	20 70-130	
Petroleum Hydi	rocarbons C6 - C35, Soil	512.187	515.849	500.000000	ND	2.1 102 0.7	20 70-130 20	
LCS	Laboratory Control Sample		GCL051906	157296-1			36/21/2006 1	938
Para	meter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hyd	rocarbons C6 - C12, Soil rocarbons C12 - C28, Soil rocarbons C6 - C35, Soil	234.799 281.051 515.849	•	250.000000 250.000000 500.000000	ND ND ND	93.9 112.4 103.2	70-130 70-130 70-130	
МВ	Method Blenk		GC061306	157296-1		1	06/21/2006 1	905
Para	meter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hyd Petroleum Hyd	rocarbons C6 - C12, Soil rocarbons C12 - C28, Soil rocarbons C28 - C35, Soil rocarbons C6 - C35, Soil	ND ND ND ND						
MS	Matrix Spike		GCS061906	318024-2		ı	06/21/2006 2	151
Para	meter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hyd	rocarbons C6 - C12, Soil rocarbons C12 - C28, Soil rocarbons C6 - C35, Soil	232.221 174.600 406.821		250.000000 250.000000 500.000000	ND 173.813 354.574	93 0 10	70-130 70-130 70-130	A A
MSD	Matrix Spike Duplicate		GCS061906	318024-2		ſ	06/21/2006 2	224
Para	meter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hyd	rocarbons C6 - C12, Soil	222.920	232.221	250.000000	ND	89	70-130	
Petroleum Hyd	rocarbons C12 - C28, Soil	184.975	174.600	250,000000	173.813	4.1	20.0 70-130	A
Petroleum Hyd	rocarbons C6 - C35, Soil	407.896	406.821	500.000000	354.574	5.8 11 0.3	20.0 70-130 20.0	A

Page 39 * %=% REC, R=RPD, A=ABS Diff., D=% Diff.



Job Number.: 318054	QUALITY	CONTROL	RESULT		t Date.: 07/10,	/2006	
CUSTOMER: ATC Associates, Inc.	PROJE	CT: 9003 N. MAI	N	ATTN:			
QC Type Description		Reag. Code	Lab	ID Dilu	tion Factor	Date Ti	me
LCD Laboratory Control Sample	Duplicate	GCL 051906	157449-1			06/23/2006 1:	516
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hydrocarbons C6 - C12, Water	337.174	356.667	333.333333	ND	101	70-130	
Petroleum Hydrocarbons C12 - C28, Water	300.856	321.956	333.333333	ND	5.6 90	20 70-130	
Petroleum Hydrocarbons C6 - C35, Water	638.030	678.622	666.666667	ND	6.8 96 6.2	20 70-130 20	
LCS Laboratory Control Sample		GCL051906	157449-1			06/23/2006 1	443
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hydrocarbons C6 - C12, Water Petroleum Hydrocarbons C12 - C28, Water Petroleum Hydrocarbons C6 - C35, Water	356.667 321.956 678.622		333.333333 333.333333 666.666667	ND	107.0 96.6 101.8	70-130 70-130 70-130	
MB Method Blank		GC961306	157449-1		1	06/23/2006 1	410
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hydrocarbons C6 - C12, Water Petroleum Hydrocarbons C12 - C28, Water Petroleum Hydrocarbons C28 - C35, Water Petroleum Hydrocarbons C6 - C35, Water	ND ND ND ND						
SB Spiked Blank		GCS061906	157449-1		(06/23/2006 1	549
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hydrocarbons C6 - C12, Water Petroleum Hydrocarbons C12 - C28, Water Petroleum Hydrocarbons C6 - C35, Water	369.036 387.017 756.053		333.333333 333.333333 666.666667		110.7 116.1 113.4	70-130 70-130 70-130	
SBD Spiked Blank Duplicate		GCS061906	157449-1			06/23/2006 1	623
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hydrocarbons C6 - C12, Water	345.353	369.036	333.333333	ND	103.6	70-130 20	
Petroleum Hydrocarbons C12 - C28, Water	316.809	387.017	333.333333	ND	6.6 95.0	70-130	
Petroleum Hydrocarbons C6 - C35, Water	662.162	756,053	666.666667	ND	20.0 99.3 13.2	20 70-130 20	



	Job Number.: 318054	QUALITY	CONTRO	L RESUL		rt Date.: 07/10,	/2006	
CUSTOMER: ATO	C Associates, Inc.	PROJEC	CT: 9003 N. MA	iIN	ATTN	:		
QC Type	Description		Reag. Coo	le Lab	ID Dil	ution Factor	Date Tir	пе
	: SW-846 8315 iption.: Formaldehyde by HI	PLC		: 157630	g/L	Analyst	: jps	
LCS	Laboratory Control Sample		LC050106	157516			06/26/2006 1	024
Parame	eter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Formaldehyde, l	Formaldehyde, Water 1069.64			10000.		107.0 39-153		
МВ	Method Blank			157516				009
Parame Formaldehyde, i	eter/Test Description Water	QC Result ND	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
мѕ	Matrix Spike		LC050206	318178-	1		06/26/2006 1	055
Param	eter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Formaldehyde, Liquid 991.05				10000.	29.92	96	39-153	
MSD	Matrix Spike Duplicate		LC050206	318178-	1		06/26/2006 1	110
Param	eter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Formaldehyde,	Liquid	904.06	991.05	10000.	29.92	87.4 9.5	70-130 20	



Method: GC Volatile Organ	cs	Metho	d Code	.: 8021		Prep Batch:
Batch(s): 157626		Test	Equipment Code: BTEX02			
ab ID DT Sample ID		Date	ATFT	ATFTB	BFB	BFBB
7626- 1 LCS 7626- 1 MB 8054- 4 B-1 8054- 5 B-2 8054- 6 B-3		06/26/2006 06/26/2006 06/26/2006 06/26/2006 06/26/2006	107.2 116.0 185.8A 117.5 117.0	108.0 115.1 142.2A 116.0 117.0	105.3 111.9 107.7 113.7 112.2	107.7 113.7 98.2 111.6 116.5
est Test Description	Limits	;				
FT a,a,a-Trifluorotoluene FTB a,a,a-Trifluorotoluene Column B BFB (Surrogate) BB BFB (Surrogate) Column B	70 - 13 B 70 - 13 64 - 13 64 - 13	5 6				
Method: GC Volatile Organ Batch(s): 157626	ics		d Code Matrix			Prep Batch: Equipment Code: BTEX02
ab ID DT Sample ID		Date	ATFT	ATFTB	BFB	BFBB
7626- 1 MB 7399- 1 MS BA1R 4' 7399- 1 MSD BA1R 4'		06/26/2006 06/26/2006 06/26/2006	119.7 114.8 114.8	115.4 115.2 115.0	111.6 109.7 111.0	114.0 111.6 112.3
est Test Description	Limits	.				
FT a,a,a-Trifluorotoluene FTB a,a,a-Trifluorotoluene Column B BFB (Surrogate) BB BFB (Surrogate) Column B	70 - 13 B 70 - 13 64 - 13	5 6				
Method: GC Volatile Organ Batch(s): 157581	ics		d Code Matrix			Prep Batch: Equipment Code: BTEX02
ab ID DT Sample ID		Date	ATFT	ATFTB	BFB	BFBB
7581- 1 LCS 7581- 1 MB 7581- 1 SB 7581- 1 SBD 8054- 1 B-1 12-14 8054- 2 B-2 20-22 8054- 3 B-3 20-22		06/23/2006 06/23/2006 06/23/2006 06/23/2006 06/23/2006 06/23/2006 06/23/2006	104.0 111.3 103.6 106.9 94.6 153.3A 86.1	104.1 113.3 106.8 108.7 96.3 183.1A 86.0	107.2 106.2 93.8 110.7 92.0 15.7A 85.0	102.3 106.0 103.4 106.9 97.5 103.4 91.3
est Test Description	Limits	1				
FT a,a,a-Trifluorotoluene FTB a,a,a-Trifluorotoluene Column B BFB (Surrogate) BB BFB (Surrogate) Column B	50 - 15 B 50 - 15 50 - 15 50 - 15	i0 i0				



SURROGATE RECOVERIES REPORT

Job Number.: 318054

Report Date.: 07/10/2006

CUSTOMER: ATC Associates, Inc.

PROJECT: 9003 N. MAIN

ATTN: Patrick Dworsczyk

M B	Method: Direct Analytical TPH Method TX 1005 Batch(s): 157402 157465				d Code: TX1005 Matrix: Soil	Prep Batch: Equipment Code:		
Lab ID		DT	Sample ID	_	Date	OTERPH		
157296-	1 LC				06/21/2006	93.34		
157296-	1 LC	S			06/21/2006	94.02		
157296-	1 MB				06/21/2006	89.50		
318024-	2 MS		EPO-45-1-(5'-6')		06/21/2006	85.43		
318024-	2 MS)	EPO-45-1-(51-61)		06/21/2006	89.00		
318054-	1		B-1 12-14		06/22/2006	95.08		
318054-	2		8-2 20-22		06/22/2006	93.28		
318054-	3		8-3 20-22		06/22/2006	91.91		
Test	Te	st Des	scription	Limits				
OTERPH	0-	Terphe	enyl	70 - 13	0			

	Method: Direct Analytical TPH Method TX 1005 Batch(s): 157591				ch: 157449 Code: EXTGC12
Lab ID		DT	Sample ID	Date OTERPH	
157449-	1 LCD			06/23/2006 96.46	
157449-	1 LCS			06/23/2006 102.3	
157449-	1 MB			06/23/2006 101.4	
	1 SB			06/23/2006 116.6	
	1 SBD			06/23/2006 104.9	
318054-	4		B-1	06/23/2006 102.8	
318054-	-		8-2	06/23/2006 101.0	
318054-			B-3	06/24/2006 107.3	
Test	Test	t Des	cription	Limits	
OTERPH	0-Te	erphe	nyl	70 - 130	



QUALITY ASSURANCE METHODS

REFERENCES AND NOTES

Report Date: 07/10/2006

REPORT COMMENTS

- All pages of this report are integral parts of the analytical data. Therefore, this report should be reproduced only in its entirety.
- 2) Reporting limits are adjusted for sample size used, dilutions and moisture content if applicable.
- 3) According to 40CFR Part 136.3, pH, Chlorine Residual, and Dissolved Oxygen analyses are to be performed immediately after aqueous sample collection. When these parameters are not indicated as field,(e.g. pH field) they were not analyzed immediately, but as soon as possible on laboratory receipt.
- 4) For all USACE projects, the QC limits are based on "mean +/- 2 sigma", which are the warning limits.

General Information:

- Cresylic Acid is the combination of o,m and p-Cresol. The combination is reported as the final result.
- m-Cresol and p-Cresol co-elute. The result of the two is reported as either m&p-cresol or as p-cresol.
- m-Xylene and p-Xylene co-elute. The result of the two is reported as m,p-Xylene.
- N-Nitrosodiphenylamine decomposes in the gas chromatograph inlet forming dipheylamine and, consequently, may be detected as diphenylamine.
- Methylene Chloride and Acetone are recognized potential laboratory contaminants. Its presence in the sample up to five times the amount reported in the blank may be attributed to laboratory contamination.
- Trimethysilyl(Diazomethane) is used to esterify acid herbicides in Method SW-846 8151A.
- For Inorganic analyses, duplicate QC limits are determined as follows: If the sample result is less than or equal to 5 times the reporting limit, the RPD limit is equal to the reporting limit. If the sample result is greater than 5 times the reporting limit, the RPD limit is the method defined RPD.
- For TRRP reports, the header on the column RL is equivalent to a MQL/PQL.

Explanation of Qualifiers:

- U This qualifier indicates that the analyte was analyzed but not detected.
- J (Organics only) This qualifier indicates that the analyte is an estimated value between the RL and the MDL.
- B (Inorganics only) This Qualifier indicates that the analyte is an estimated value between the RL and the MDL.
- N (Organics only) This flag indicates presumptive evidence of a compound. This flag is only used for tentatively identified compounds (TICs), where the identification is based on a mass spectral library search. It is applied to all TIC results. For generic charachterization of a TIC, such as "chlorinated hydrocarbon", the "N" flag is not used.

Explanation of General QC Outliers:

- A Matrix interference present in sample.
- a MS/MSD analyses yielded comparable poor recoveries, indicating a possible matrix interference. Method performance is demonstrated by acceptable LCS recoveries.
- b Target analyte was found in the method blank.
- M QC sample analysis yielded recoveries outside QC acceptance criteria. This sample was reanalyzed.
- L LCS analysis yielded high recoveries, indicating a potential high bias. No target analytes were observed above the RL in the associated samples.
- G Marginal outlier within 1% of acceptance criteria.
- r RPD value is outside method acceptance criteria.
- C Poor RPD values observed due to the non-homogenous nature of the sample.
- O Sample required dilution due to matrix interference.
- D Sample reported from a dilution.
- d Spike and/or surrogate diluted.
- P The recovery of this analyte is outside default QC limits. The data is accepted and will be used to calculate in-house statistical limits.
- E The reported concentration exceeds the instrument calibration.
- F The analyte is outside QC limits. The sample data is accepted since this analyte is not reported in associated samples.
- H Continuing Calibration Verification (CCV) standard is not associated with the samples reported.



QUALITY ASSURANCE METHODS REFERENCES AND NOTES

Report Date: 07/10/2006

q - See the subcontract final report for qualifier explanation.

- W The MS/MSD recoveries are outside QC acceptance criteria because the amount spiked is much less than the amount found in the sample.
- K High recovery will not affect the quality of reported results.

Z - See case narrative.

Explanation of Organic QC Outliers:

- e Method blank analysis yielded phthalate concentrations above the RL. Phthlates are recognized potential laboratory contaminants. Its presence in the sample up to five times the amount reported in the blank may be attributed to laboratory contamination.
- S Sample reanalyzed/reextracted due to poor surrogate recovery. Reanalysis confirmed original analysis indicating a possible matrix interference.

T - Sample analysis yielded poor surrogate recovery.

- R The RPD between the two GC columns is greater than 40% and no anomalies are present. The higher result is reported as per EPA Method 8000B.
- I The RPD between the two GC columns is greater than 40% and anomalies are present. The lower of the two results has been reported.

X - Gaseous compound. In-house QC limits are advisory.

- Ketone compounds have poor purge efficiency. In-house QC limits are advisory.

f - Surrogate not associated with reported analytes.

Explanation of Inorganic QC Outliers:

- Q Method blank analysis yielded target analytes above the RL. Associated sample results are greater than 10 times the concentrations observed in the method blank.
- The RPD control limit for sample results less than 5 times the RL is +/- the RL value. Sample and duplicate results are within method acceptance criteria.

e - Serial dilution failed due to matrix interference.

- g Sample result quantitated by Method of Standard Additions (MSA) due to the analytical spike recovery being below 85 percent. The correlation coefficent for the MSA is greater than or equal to 0.995.
- BOD/cBOD seed value is not within method acceptance criteria. Due to the nature of the test method, the sample cannot be reanalyzed.
- 1 BOD/CBOD LCS value is not within method acceptance criteria. Due to the nature of the test method, sample cannot be reanalyzed.

- Spiked sample recovery is not within control limits.

n - Sample result quantitated by Method of Standard Additions (MSA) due to the analytical spike recovery being below 85 percent. The correlation coefficient for the MSA is less than 0.995.

* - Duplicate analysis is not within control limits.

Abbreviations:

- Designation given to identify a specific extraction, digestion, preparation, or analysis set. Batch

- Continuing Calibration Verification CCV

- Low level standard check - GFAA, Mercury CRA

- Low level standard check - ICP CR I

Dil Fac - Dilution Factor - Secondary dilution analysis

- Detection Limit Factor DLFac

- Duplicate DU

- Extraction Blank (TCLP, SPLP, etc.)

EB - Initial Calibration ICAL

ICB - Initial Calibration Blank

ICV Initial Calibration Verification

Interference Check Sample A - ICP ISA ISR

- Interference Check Sample B - ICP - Laboratory Control Duplicate LCD

LCS - Laboratory Control Sample



QUALITY ASSURANCE METHODS REFERENCES AND NOTES

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MB - Method Blank - Method Duplicate MD - Method Detection Limit MDI - Method Quantitation Limit (TRRP) MQL - Matrix Spike - Matrix Spike Duplicate MSD - Not Detected ND - Preparation Blank PR PREPF - Preparation Factor - Reporting Limit RPD - Relative Percent Difference - Relative Response Factor RRF - Retention Time RT - Sample Quantitation Limit (TRRP) SQL TIC - Tentatively Identified Compound

Method References:

- (1) EPA 600/4-79-020 Methods for the Analysis of Water and Wastes, March 1983.
- (2) EPA 600/R-94-111 Methods for the Determination of MEtals in Environmental Samples, Supplement I, May 1994.
- (3) EPA SW846 Test Methods for Evaluating Solid Waste, Third Edition, September 1986; Update I July 1992; Update II, September 1994, Update IIA August 1993; Update IIB, January 1995; Update III, December 1996, Update IVA January 1998, Update IVB November 2000.
- (4) Standard Methods for the Examination of Water and Wastewater, 16th Edition (1985), 17th Edition (1989), 18th Edition (1992), 19th Edition (1995), 20th Edition (1998). (5) HACH Water Analysis Handbook 3rd Edition (1997).

- Federal Register, July 1, 1990 (40 CFR Part 136 Appendix A).

 Compendium of Methods for the Determination of Toxic Organic Compounds in Ambient Air, 2nd Edition, (7) January 1997.
- (9) Diagnosis and Improvement of Saline and Alkali Soils, Agriculture Handbook No. 60, United States Department of Agriculture, 1954.



LABORATORY CHRONICLE

Job Number: 318054

Date: 07/10/2006

CUSTOMER: ATC Ass	ociates, inc. PROJE	CT: 9003 N	. MAIN		A	TTN: Patrick D	lworaczy	k
ab ID: 318054-1	Client ID: B-1 12-14	Date Re	cvd: 06/	21/2006	Sample	Date: 06/21/20	006	
METHOD	DESCRIPTION			PREP BT		DATE/TIME AN		DILUTIO
METHOD	Data Package Validation	1	158535		"(")	07/10/2006	0000	0.20.70
THROC 100E	Direct Analytical TPH Method TX 1005	i		157296		06/22/2006	2038	1.0000
TNRCC 1005	Direct Analytical Irm Method IA 1005	i	157594	131270		06/26/2006	1045	,,,,,,,,
	Extractable GC Data Package Production						0800	
SW846 8315	Extraction Formaldehyde Waters	1	157516	457547		06/23/2006		4 000
sw-846 8315	Formaldehyde by HPLC	1		157516		06/26/2006	1156	1.000
SW-846 8021B	GC Volatile Organics	1	157581			06/23/2006	2050	1.0000
	GC Volatiles Data Package Production	1						
SM-2540 G Mod.	Moisture (Total + Fixed Solids, Ash)	1	157440			06/22/2006	1630	
TNRCC TX-1005	TNRCC 1005 Extraction (Ultrasonic)	1	157296			06/21/2006	1600	
SW-846 1311	Toxicity Characteristic Leachate Proced.	1	157473			06/22/2006	2230	
ab ID: 318054-2	Client ID: B-2 20-22	Date Re	cvd: 06/	21/2006	Sample	Date: 06/21/20	006	
METHOD	DESCRIPTION			PREP BT		DATE/TIME AN		DILUTIO
	Direct Analytical TPH Method TX 1005	1		157296	w(u/	06/22/2006	2144	1.0000
TNRCC 1005				131290				1.0000
	Extractable GC Data Package Production	1	157594			06/26/2006	1045	
SW846 8315	Extraction Formaldehyde Waters	1	157516			06/23/2006	0800	
SW-846 8315	Formaldehyde by HPLC	1		157516		06/26/2006	1212	1.000
SW-846 80218	GC Volatile Organics	1	157581			06/23/2006	2130	1.0000
SM-2540 G Mod.		1	157440			06/22/2006	1630	
TNRCC TX-1005	TNRCC 1005 Extraction (Ultrasonic)	1	157296			06/21/2006	1600	
SW-846 1311	Toxicity Characteristic Leachate Proced.	i	157473			06/22/2006	2230	
		D		24 /2004	Comple	Data 06/21/2	204	
ab ID: 318054-3	Client ID: B-3 20-22		cvd: 06/			Date: 06/21/20		
METHOD	DESCRIPTION			PREP BT	#(S)	DATE/TIME A		DILUTI
TNRCC 1005	Direct Analytical TPH Method TX 1005	1		157296		06/22/2006	2217	1.0000
	Extractable GC Data Package Production	1	157594			06/26/2006	1045	
SW846 8315	Extraction Formaldehyde Waters	1	157516			06/23/2006	0800	
SW-846 8315	Formaldehyde by HPLC	1	157630	157516		06/26/2006	1227	1.000
SW-846 8021B	GC Volatile Organics	1	157581			06/23/2006	2230	1.0000
		i	157440			06/22/2006	1630	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
SM-2540 G Mod.		j	157296				1600	
TNRCC TX-1005	TNRCC 1005 Extraction (Ultrasonic)	•				06/21/2006		
sw-846 1311	Toxicity Characteristic Leachate Proced.	1	157473			06/22/2006	2230	
ab ID: 318054-4	Client ID: B-1		cvd: 06/			Date: 06/21/20		
METHOD	DESCRIPTION			PREP BT	#(S)	DATE/TIME AN		DILUTIO
TNRCC 1005	Direct Analytical TPH Method TX 1005	1	157591	157449		06/23/2006	2303	1.0000
SW846 8315	Extraction Formaldehyde Waters	1	157516			06/23/2006	0800	
SW-846 8315	Formaldehyde by HPLC	1	157630	157516		06/26/2006	1258	1.000
SW-846 8021B	GC Volatile Organics	1	157626			06/26/2006	1020	1.0000
TNRCC TX-1005	TX-1005 Extraction Water	i	157449			06/22/2006	1430	,,,,,,,,,
-1- rh - 7400F/ F	nii In- n-3	Doto Do	cvd: 06/	21 (2004	Comple	Datas 04/21/20	204	
ab ID: 318054-5	Client ID: B-2					Date: 06/21/20		DYLLETY
METHOD	DESCRIPTION			PREP BT	#(5)	DATE/TIME AN		DILUTIO
TNRCC 1005	Direct Analytical TPH Method TX 1005	1		157449		06/23/2006	2336	1.0000
SW846 8315	Extraction Formaldehyde Waters	1	157516			06/23/2006	0800	
SW-846 8315	Formaldehyde by HPLC	1	157630	157516		06/26/2006	1313	1.000
SW-846 8021B	GC Volatile Organics	1	157626			06/26/2006	1040	1.0000
TNRCC TX-1005	TX-1005 Extraction Water	1	157449			06/22/2006	1430	
ab ID: 318054-6	Client ID: B-3	Date Pe	cvd: 06/	21/2006	Sample	Date: 06/21/20	006	
								DILUTI
	DESCRIPTION	_		PREP BT	かしごと	DATE/TIME AN		
METHOD	Direct Analytical TPH Method TX 1005	1	12/27	157449		06/24/2006	0009	1.0000
TNRCC 1005						A	***	
	Extraction Formaldehyde Waters	1	157516			06/23/2006	0800	
TNRCC 1005		1		157516		06/23/2006 06/26/2006	0800 1329	1.000



LABORATORY CHRONICLE

Job Number: 318054

Date: 07/10/2006

CUSTOMER: ATC Associates, Inc.

PROJECT: 9003 N. MAIN

ATTN: Patrick Dworaczyk

Lab ID: 318054-6 METHOD

Client ID: B-3
DESCRIPTION

TNRCC TX-1005 TX-1005 Extraction Water

Date Recvd: 06/21/2006 Sample Date: 06/21/2006

RUN# BATCH# PREP BT #(S) 157449

DATE/TIME ANALYZED DILUTION

06/22/2006 1430



LIMITED PHASE II ENVIRONMENTAL SITE ASSESSMENT

CITY OF HOUSTON 9003 NORTH MAIN STREET HOUSTON, TX ATC PROJECT NO. 73.17331.0074

June 30, 2006

Prepared by:

ATC Associates Inc. 3928 Bluebonnet Drive Stafford, Texas 77477 Telephone: (281) 240-0154

Facsimile: (281) 240-8909

Prepared for:

The City of Houston 900 Bagby, 2nd Floor Houston, TX 77002 Telephone: (713) 247-3232

Facsimile: (713)

Patrick Dworaczyk, P.G., CAPM Project Manager

Mark Smith, P.G., CAPM Senior Geologist



ASBESTOS AND LEAD-BASED PAINT SURVEY

City of Houston 9003 North Main Street Houston, Texas

ATC Project Number 73.17331.0074

June 2006

PREPARED FOR:

City of Houston P.O. Box 1562 Houston, Texas 77251-1562 Contract No. C50597

PREPARED BY:

ATC Associates Inc. 3928 Bluebonnet Drive Stafford, Texas 77477



3928 Bluebonnet Drive Stafford, Texas 77477 www.atcassociates.com Office 281-240-0154 Fax 281-240-8909

June 29, 2006

Mr. Gabriel Mussio
Division Manager
Energy and Environmental Management Division
City of Houston
Building Services Department
900 Bagby, 2nd Floor
Houston, TX 77002

Re: Asbestos and Lead-Based Paint Survey
City of Houston
9003 North Main Street, Houston, Texas

ATC Project No. 73.17331.0074

Contract No. C50597

Dear Mr. Mussio:

Enclosed is the asbestos and lead-based paint survey of the City of Houston property located at 9003 North Main Street in Houston, Texas.

Ms. Jennifer Boone of ATC performed the survey on June 28, 2006. The enclosed report describes the results of sampling and analysis and provides a homogeneous area report. All original laboratory reports and ATC's submittals are included as appendices.

We appreciate the opportunity to provide environmental consulting services to the City of Houston and look forward to assisting you with future consulting services. If you have any questions or need additional assistance, please call 281.240.0154.

Sincerely,

ATC Associates Inc.

Jennifer L. Boone

Sr. Environmental Scientist

TDSHS IAC Lic. #10-5554

migke Boone

Catherine G. McLain

Industrial Hygiene Department Manager

TDSHS IAC Lic. #10-5451

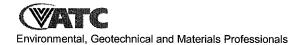


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EXECUTIVE SUMMARY

ATC Associates Inc. (ATC) performed an asbestos and lead-based paint survey of the City of Houston property located at 9003 North Main Street in Houston, Texas. The scope of work included: review of previous documentation (if provided), inspecting and sampling for suspect Asbestos-Containing Building Materials (ACBM) and Lead-Based Paint Materials (LBPM), and identifying, quantifying and assessing confirmed ACBM and LBPM.

The vacant facility was formerly used as an office and storage facility by the Street Maintenance Department. The site contains a one-story office building with an attached warehouse area in the eastern section of the property, a metal warehouse building at the north end of the property, and a canopy covered truck wash area on the southwestern portion of the site. Building exteriors are generally metal siding except for three brick walls on the office building that face North Main Street. The buildings have corrugated metal roofs and concrete slab foundations. Interior finishes are composed of concrete block, plywood and wallboard walls, textured wallboard walls, 1' x 1' suspended ceiling tiles, 2' x 2' suspended ceiling tiles, 2' x 4' suspended ceiling tiles, 12" x 12" floor tiles, concrete floors and carpet. No interior suspect asbestos-containing materials were found in the Car Wash Building or the Warehouse Building.

ATC's Texas Department of State Health Services (TDSHS) Inspector Ms. Jennifer Boone (Asbestos License No. 10-5554, performed the survey on June 28, 2006. The exterior and roof of the buildings were not included in the scope of work.

Asbestos Survey

Asbestos-containing material (ACM) is defined by the <u>Texas Asbestos Health Protection Rules</u> (TAHPR), March 2003, as materials or products that contain more than 1.0% of any kind or combination of asbestos, as determined by the Environmental Protection Agency (EPA) recommended methods as listed in <u>EPA/600/R-93/116</u>, July 1993 "Method for the Determination of Asbestos in Bulk Building Materials". This means any one material component of a structure or any layer of a material sample.

Using the guidelines set by TDSHS, samples of suspect ACBM were collected and sent under chain-of-custody to Hygeia Laboratories Inc. in Miami, Florida (TDSHS License No. 30-0230). The analytical results are summarized in the following report. Table 1, Summary of Suspected ACBM Bulk Samples lists materials sampled, sample locations, and sample results. Table 2, Summary of ACBM Assessment, lists confirmed ACBM, their location, estimated quantity, and hazard assessment.

Based on the results presented in Table 1, the following materials were identified through laboratory analysis as containing greater than 1% asbestos:

 Approximately 232 linear feet of gray interior window glazing located in Offices 6, 7, 8, and Restroom A.



Based on the results of the limited survey, ATC proposes the following options to the City of Houston:

- Prior to renovation, ACBM with the potential for disturbance must be properly abated and disposed of in compliance with the TDSHS TAHPR and National Standards for Hazardous Air Pollutants (NESHAP).
- If additional suspect ACBM other than those identified in this report are observed during renovation activities, appropriate samples should be collected and analyzed for asbestos content prior to disturbance.
- Prior to any demolition activities a thorough survey of roofing and exterior materials should be completed.

Limited Lead-Based Paint Survey

Lead-Based Paint (LBP) is defined by the <u>HUD Guidelines for the Evaluation and Control of Lead-Based Paint Hazards in Housing</u>, June 1995, revised 1997 as any paint, varnish, shellac, or other coating that contains lead equal to or greater than 1.0 milligrams per square centimeter (mg/cm²) as measured by XRF or laboratory analysis; or in excess of 0.5 percent by weight (5,000 ppm) as measured by laboratory analysis. However, it should be noted that the Occupational Safety and Health Act (OSHA) lead construction standard, 29 CFR §1926.62, regulates workers involved in any activity whereby lead-containing materials could be disturbed resulting in airborne lead exposure. Therefore, the City of Houston standard calls for any material containing lead in amounts greater than 0.0 mg/kg requires an OSHA exposure assessment for the possible exposure to lead hazards.

Using the guidelines set by HUD, a protocol was developed for the limited lead-based paint survey. The survey began with a visual inspection and assessment of the condition of suspect LBPM. Paint chips were collected from materials randomly selected based on color, substrate and paint history. The collected samples were submitted for analysis to Hygeia Laboratories Inc. in New York, New York (AIHA Accreditation No. 100229).

Analytical results did show the presence of one painted materials meeting the HUD definition of LBPM in the tested locations. This material includes:

• Approximately 2,000 square feet of red primer paint on the structural steel in the Office Building which contains 0.66% lead by weight.

Analytical results showed one painted material identified as containing detectable concentrations of lead and is therefore subject to the OSHA requirements. This material includes:

• Approximately 5,000 square feet of off-white paint on the interior plywood walls and wood stairway in the warehouse area of the Office Building which contains 0.02% lead by weight.



Based on the analytical results, ATC makes the following recommendations:

- It is recommended contractors working with any painted materials at this site maintain compliance with the OSHA lead construction standard, 29 CFR §1926.2, during any construction and renovation activities impacting these materials. The OSHA lead construction standard, 29 CFR §1926.62, regulates workers involving any activity whereby lead-containing materials could be disturbed resulting in airborne lead exposure.
- Contractors working with the known LBPM must maintain compliance with the OSHA lead construction standard, 29 CFR 1926.62, during any construction and renovation activities impacting these materials.
- LBPM may be abated prior to demolition or renovation to ensure an environmentally safe work area, metal components removed and sent to a smelter that accepts lead containing materials, or encapsulated with layers of lead-free paint and monitored under an operations and maintenance plan.
- For demolition and disposal purposes under the federal Resource Conservation and Recovery Act (RCRA) the LBPM should be tested using toxicity characteristic leaching procedure (TCLP) to determine the hazardous waste classification.

1.0 OBSERVATIONS

The vacant facility was formerly used as an office and storage facility by the Street Maintenance Department. The site contains a one-story office building with an attached warehouse area in the eastern section of the property, a metal warehouse building at the north end of the property, and a canopy covered truck wash area on the southwestern portion of the site. Building exteriors are generally metal siding except for three brick walls on the office building that face North Main Street. The buildings have corrugated metal roofs and concrete slab foundations. Interior finishes are composed of concrete block, plywood and wallboard walls, textured wallboard walls, 1' x 1' suspended ceiling tiles, 2' x 2' suspended ceiling tiles, 2' x 4' suspended ceiling tiles, 12" x 12" floor tiles, concrete floors and carpet. No interior suspect asbestos-containing materials were found in the Car Wash Building or the Warehouse Building.

2.0 STANDARDS AND GUIDELINES

Below is a brief summary of applicable federal and state regulations for asbestos in building materials.

2.1 OSHA Asbestos in Construction Standard at 29 CFR § 1926.1101

The OSHA Asbestos Standard regulates workers in construction, demolition, and maintenance who may be occupationally exposed to asbestos-containing products. Asbestos-related construction work involves any demolition or salvage of structures where asbestos is present; removal or encapsulation of materials containing asbestos; construction, alteration, repair, maintenance, or renovation of structures, substrates, or portions thereof, that contain asbestos; installation of products containing asbestos; asbestos spill/emergency cleanup; and transportation, disposal, storage, containment of and housekeeping activities involving asbestos or products containing asbestos, on the site or location at which construction activities are performed whereby the asbestos-containing material could be disturbed resulting in asbestos exposure.

The standard classifies different activities involving asbestos based on their potential for disturbance. It also states the employer shall sample the air in the worker's breathing zone to determine asbestos exposure. Permissible Exposure Limits (PEL) and Excursion Limits are established in the OSHA standard for comparison. The employer shall ensure that no employee is exposed to an airborne concentration of asbestos in excess of 0.1 fiber per cubic centimeter of air as an eight hour time-weighted average (TWA) or an airborne concentration of 1.0 fiber per cubic centimeter of air (1 f/cc) as averaged over a sampling period of 30 minutes. All employers of employees exposed to asbestos hazards must comply with applicable protective provisions to protect their employees. The standard designates approved work practices for dealing with asbestos, establishes air monitoring requirements, and sets requirements for appropriate respiratory protection.

2.2 EPA National Emission Standards for Hazardous Air Pollutants (NESHAP) 40 CFR Part 61

The NESHAP rules set a ten-day notification requirement for planned demolitions and renovations. Additionally, recordkeeping requirements were established for waste disposal. The rules categorize ACM based on the potential to release fibers.

2.3 Texas Asbestos Health Protection Rules (TAHPR) Title 25, Part 1, §295

ACM is defined by the <u>Texas Asbestos Health Protection Rules</u> (TAHPR), March 2003, as materials or products that contain more than 1.0% of any kind or combination of asbestos, as determined by the EPA recommended methods as listed in <u>EPA/600/R-93/116</u>, July 1993 "Method for the Determination of Asbestos in Bulk Building <u>Materials</u>". This means any one material component of a structure or any layer of a material sample.

The TAHPR address the problem of limiting the exposure of an individual to asbestos fibers by regulating asbestos disturbance activities in buildings that afford public access or occupancy and in commercial buildings. These regulated activities apply to all persons disturbing, removing, encapsulating, or enclosing asbestos within public buildings for any purpose, including repair, renovation, dismantling, demolition, installations, or maintenance operations, or any activity that may involve the disturbance or removal of ACM whether intentional or unintentional. Also included are the qualifications for licensure of persons and requirements for compliance with these sections and all applicable standards of the United States Environmental Protection Agency and the United States Occupational Safety and Health Administration as adopted.

Below is a brief summary of applicable federal and state regulations for lead in paint.

2.4 United States Housing and Urban Development (HUD)

LBP is defined by the <u>HUD Guidelines for the Evaluation and Control of Lead-Based Paint Hazards in Housing</u>, June 1995, revised 1997 as any paint, varnish, shellac, or other coating that contains lead equal to or greater than 1.0 mg/cm² as measured by XRF or laboratory analysis; or in excess of 0.5 percent by weight (5,000 ppm, 5,000 mg/kg, or 5,000 μg/g) as measured by laboratory analysis.

2.5 OSHA Lead in Construction Standard at 29 CFR § 1926.62

The OSHA Lead Standard regulates workers in construction, demolition, and maintenance who may be occupationally exposed to lead-containing products. Lead-related construction work involves any construction, repair, painting, demolition, renovation, removal or encapsulation, alteration, installation of lead products, emergency cleanup, transportation, disposal, storage, containment, and maintenance work whereby the lead-containing material could be disturbed resulting in lead exposure.

The standard requires all employers to provide an exposure assessment for the possible exposure to lead hazards. One component of the mandatory exposure assessment involves sampling the air in the worker's breathing zone to determine lead exposure. Action Levels and Permissible Exposure Limits (AL and PEL) are established in the OSHA standard for comparison. All employers who may expose workers above the PEL of 50 micrograms per cubic meter of air ($\mu g/m^3$) averaged over an eight hour period must develop a written compliance program prior to the start of each job. The standard addresses the circumstances under which employees must wear personal protective equipment. Employers must make available medical exams for workers as well as testing for blood lead levels.

Regarding tasks involving lead-containing materials, the employer must perform an employee exposure assessment and document that the employee performing the task is not exposed above the AL of 30 μ g/m³, or the employer shall treat the employee as if the employee were exposed above the PEL (50 μ g/m³).

3.0 SAMPLING PROCEDURES

Suspected ACBM

П

The inspector followed the sampling procedure in accordance with TDSHS TAHPR.

Suspect materials, which are alike in appearance and application, were sampled as a homogeneous area. Suspect homogeneous areas are divided into three classifications: 1) Surfacing materials: spray-applied or troweled on material, 2) Thermal Systems Insulation: pipe, boiler, tank, or flue insulation, and 3) Miscellaneous: other suspect material, including floor tile, floor tile mastic, sheet vinyl flooring, ceiling tile and panel, insulation mastics, cove base mastic, window caulking/glazing, and exterior siding (transite).

Suspect materials sampled and analyzed should be considered homogeneous area if: 1) They exhibit similar physical characteristics, and 2) The application of the sampled material can be correlated to the application of unsampled material.

Representative sampling was based on: 1) Distribution of the suspect materials throughout the homogeneous area, 2) The suspect material's physical characteristics and application, and 3) Random sampling patterns determined for each homogeneous area.

The asbestos survey included the collection of thirty-four (34) bulk samples of suspect ACBM. Where more than one sample of a homogeneous area was collected, analysis was conducted on a "stop at first positive" basis. Ms. Jennifer Boone of ATC performed the field survey on June 28, 2006. The materials sampled, sample locations, and analytical results are included in Table 1, Summary of Suspected ACBM Bulk Samples, with the bulk sample analysis report included as Appendix B.

Suspect materials sampled during our survey included the following: wall texture and wallboard joint compound, wallboard walls, 1' x 1' suspended ceiling tiles and mastic, 2' x 2' suspended ceiling tiles, 2' x 4' suspended ceiling tiles, 12" x 12" floor tiles and carpet mastic. No interior suspect asbestos-containing materials were found in the Car Wash Building or the Warehouse Building.

Suspected LBPM

The limited lead-based paint survey began with a visual inspection and assessment of the condition of suspect LBPM. Eight (8) paint chip samples were collected from materials randomly selected based on color, substrate and paint history. The collected samples were submitted for analysis to Hygeia Laboratories Inc. in New York, New York American Industrial Hygiene Association (AIHA) Accreditation No. 100229. The materials sampled, sample location, and analytical results are included in Table 3, Summary of Analysis of Paint for Lead Determination, with the bulk sample analysis report included as Appendix C.

4.0 ANALYTICAL PROCEDURES

Suspected ACBM

Collected suspected ACBM bulk samples were analyzed under polarized light microscopy (PLM) by Hygeia Laboratories Inc. in Miami, Florida, utilizing the Environmental Protection Agency's Method for the Detection of Asbestos in Bulk Insulation Samples, (EPA 600/R-93 July 1993), and the McCrone Research Institute's The Asbestos Particle Atlas as method references. ATC's laboratory is accredited by the National Voluntary Laboratory Accreditation Program (NVLAP), participates in the NVLAP and AIHA Bulk Asbestos Sample Quality Assurance Programs, and is licensed to analyze bulk asbestos samples collected in the State of Texas.

Suspected LBPM

Collected paint chips were analyzed by Hygeia Laboratories Inc. in New York, New York, utilizing the Environmental Protection Agency's Analytical Method 7420 and Digestion Method 3050. ATC's laboratory is accredited by the AIHA and certified under the Environmental Lead Laboratory Accreditation Program (ELLAP).

5.0 ANALYTICAL RESULTS

Suspected ACBM

Based on the results presented in Table 1, the following materials indicated the presence of asbestos in amounts greater than 1%:

 Approximately 232 linear feet of gray interior window glazing located in Offices 6, 7, 8, and Restroom A.

Due to the sometimes difficulty in analyzing non-friable or resinously bound materials, Hygeia Laboratories recommends that these types of materials, which were not found to contain asbestos, be analyzed using alternative methods of identification, such as Transmission Electron Microscopy.

The analytical results can be found in the attached Appendix C. Table 1, Summary of Analysis of Suspected ACBM Bulk Samples lists materials sampled, sample locations, and sample results. Table 2, Summary of ACBM Assessment lists confirmed ACBM, their locations, estimated quantities, and hazard assessment.

Suspected LBPM

Based on the results presented in Table 3, the following materials indicated the presence of lead in amounts greater than 0.06%:

• Approximately 2,000 square feet of red primer paint on the structural steel in the Office Building contains 0.66% lead by weight.

The following materials indicated the presence of lead in amounts less than 0.06%:

• Approximately 5,000 square feet of off-white paint on the interior plywood walls and wood stairway in the warehouse area of the Office Building contains 0.02% lead by weight.

The analytical results can be found in the attached Appendix C. Table 3, Summary of Analysis of Paint for Lead Determination lists materials sampled, sample locations, and sample results. Table 4, Summary of LBPM Assessment lists confirmed LBPM, their locations, estimated quantities, and hazard assessment.

6.0 HAZARD CONDITION ASSESSMENT

A condition assessment refers to the process where a material's potential to release fibers or dust particles into the air is evaluated. Fibers or dust may be released inadvertently by localized disturbance, as part of a material's aging process, or when acted upon by other factors such as air

movement, impact, or vibration. Assessing a material's potential for fiber or dust release (therefore, its associated hazard risk) is accomplished by evaluating associated factors.

The hazard condition assessments given are based on the City of Houston Asbestos Hazard Categorization (AHC) and Lead Hazard Categorization (LHC) lists. The hazard associated with any ACBM or LBPM may become more extensive over time. Each building use has the potential to contribute to a change in the potential health hazard. Tables 2 and 4 provide a hazard assessment for identified ACBM and LBPM in the buildings located at 9003 North Main Street in Houston, Texas.

7.0 QUANTITY ESTIMATES

The following materials indicated the presence of asbestos in amounts greater than 1%:

• Approximately 232 linear feet of gray interior window glazing located in Offices 6, 7, 8, and Restroom A.

Analytical results did not show the presence of painted materials meeting the HUD definition of LBPM in any of the tested locations. However, one paint material was identified as containing detectable concentrations of lead and is therefore subject to the OSHA requirements. This material includes:

- Approximately 2,000 square feet of red primer paint on the structural steel in the Office Building contains 0.66% lead by weight.
- Approximately 5,000 square feet of off-white paint on the interior plywood walls and wood stairway in the warehouse area of the Office Building contains 0.02% lead by weight.

8.0 CONCLUSIONS AND RECOMMENDATIONS

Based on the results of the survey, ATC proposes the following options to the City of Houston:

- Prior to renovation, ACBM with the potential for disturbance must be properly abated and disposed of in compliance with the TDSHS TAHPR and NESHAP.
- If additional suspect ACBM other than those identified in this report are observed during renovation activities, appropriate samples should be collected and analyzed for asbestos content prior to disturbance.
- Prior to any demolition activities a thorough survey of roofing materials and inaccessible materials should be completed.

It should be noted that the EPA has not prohibited the manufacture of non-friable asbestos-containing building materials, such as vinyl floorings, mastics, and roofing materials. As a result, any future replacement materials should be checked for the presence of asbestos.

Based on the analytical results for the collected paint chip samples, ATC makes the following recommendations:

- It is recommended contractors that work with any painted materials at this site maintain compliance with the OSHA lead construction standard, 29 CFR §1926.2, during any construction and renovation activities impacting these materials. OSHA lead construction standard, 29 CFR §1926.62, regulates workers involving any activity whereby lead-containing materials could be disturbed resulting in airborne lead exposure. Therefore, any material containing lead in amounts greater than 0.0 mg/cm² should require an exposure assessment for the possible exposure to lead hazards.
- Contractors working with the known LBPM must maintain compliance with the OSHA lead construction standard, 29 CFR 1926.62, during any construction and renovation activities impacting these materials.
- LBPM may be abated prior to demolition or renovation to ensure an environmentally safe work area, metal components removed and sent to a smelter that accepts lead containing materials, or encapsulated with layers of lead-free paint and monitored under an operations and maintenance plan.
- For demolition and disposal purposes under the federal Resource Conservation and Recovery Act (RCRA) the LBPM should be tested using toxicity characteristic leaching procedure (TCLP) to determine the hazardous waste classification.

9.0 LIMITATIONS

This report has been prepared to assist the City of Houston in evaluating the ACBM and LBPM in the City of Houston facility located at 9003 North Main Street in Houston, Texas. Our objective was to perform our work with care, exercising the customary skill and competence of consulting professionals in the relevant disciplines in this region. The conclusions presented in this report are professional opinions based solely upon visual observations of the site, at the time of our investigation, and the results of laboratory analysis. The opinions presented herein apply to site conditions existing at the time of our investigation and those reasonably foreseeable. Quantity estimates of confirmed ACBM and LBPM are preliminary, based on observations made during our survey and should not be used to prepare a removal cost estimate. ATC cannot act as insurers, and no expressed or implied representation or warrant is included or intended in our report except that our work was performed, within the limits prescribed by our clients, with the customary thoroughness and competence of our profession at the time and place the services were rendered. Unsampled ACBM and LBPM may be located within walls, ceiling cavities, below flooring or grade, and other non-accessible areas. Precaution should be used in relation to these unsampled materials until their asbestos content has been determined by proper sampling and analysis. The condition of the ACBM and LBPM may change gradually or suddenly, depending upon use, maintenance or accident.

This report is intended for the sole use of the City of Houston. The scope of services performed in execution of this evaluation may not be appropriate to satisfy the needs of other users, and use or re-use of this document or the findings, conclusions, or recommendations, is at risk of said user.

ATC Associates Inc.

Jennifer L. Boone

Sr. Environmental Scientist

TDSHS IAC Lic. #10-5554

Catherine G. McLain

Industrial Hygiene Department Manager

TDSHS IAC Lic. #10-5451

TABLE 1 SUMMARY OF SUSPECTED ACBM BULK SAMPLES Office Building, 9003 North Main Street, Houston, Texas

SAMPLE NO.	DESCRIPTION/LOCATION	RESULT
B-01	Wallboard Throughout	None Detected
B-01 JC	Joint compound Throughout	None Detected
B-02	Wallboard Throughout	None Detected
B-02 JC	Joint compound Throughout	None Detected
B-03	Wallboard Throughout	None Detected
B-03 JC	Joint compound Throughout	None Detected
B-04	Wallboard Throughout	None Detected
B-04 JC	Joint compound Throughout	None Detected
B-05	Wallboard Throughout	None Detected
B-05 JC	Joint compound Throughout	None Detected
B-06	Wallboard Throughout	None Detected
B-06 JC	Joint compound Throughout	None Detected
B-07	Wallboard Throughout	None Detected
B-07 JC	Joint Compound	None Detected
B-08	Cove Base Mastic Throughout	None Detected
B-09	Cove Base Mastic Throughout	None Detected
B-10	Cove Base Mastic Throughout	None Detected
B-11	Carpet Mastic Throughout	None Detected
B-12	Carpet Mastic Throughout	None Detected
B-13	Carpet Mastic Throughout	None Detected
B-14	1' x 1' ceiling tile Offices 1 through 5 and Hallway 1	None Detected

TABLE 1 SUMMARY OF SUSPECTED ACBM BULK SAMPLES Office Building, 9003 North Main Street, Houston, Texas

SAMPLE NO.	DESCRIPTION/LOCATION	RESULT
B-14 Mastic	1' x 1' ceiling tile Mastic Offices 1 through 5 and Hallway 1	None Detected
B-15	1' x 1' ceiling tile Offices 1 through 5 and Hallway 1	None Detected
B-15 Mastic	1' x 1' ceiling tile Mastic Offices 1 through 5 and Hallway 1	None Detected
B-16	1' x 1' ceiling tile Offices 1 through 5 and Hallway 1	None Detected
B-16 Mastic	1' x 1' ceiling tile Mastic Offices 1 through 5 and Hallway 1	None Detected
B-17	12" x 12" white with tan splotches floor tile Entry Hall	None Detected
B-17 Mastic	Yellow mastic Entry Hall	None Detected
B-18	12" x 12" white with tan splotches floor tile Entry Hall	None Detected
B-18 Mastic	Yellow mastic Entry Hall	None Detected
B-19	12" x 12" white with tan splotches floor tile Entry Hall	None Detected
B-19 Mastic	Yellow mastic Entry Hall	None Detected
B-20	2' x 2' suspended ceiling tile Hallway 2	None Detected
B-21	2' x 2' suspended ceiling tile Hallway 2	None Detected
B-22	2' x 2' suspended ceiling tile Hallway 2	None Detected
B-23	12" x 12" black floor tile Hallway 2, Offices 1 through 6, Cafeteria Room, West Restrooms	None Detected
B-23 Mastic	Yellow mastic Hallway 2, Offices 1 through 6, Cafeteria Room, West Restrooms	None Detected
B-24	12" x 12" black floor tile Hallway 2, Offices 1 through 6, Cafeteria Room, West Restrooms	None Detected
B-24 Mastic	Yellow mastic Hallway 2, Offices 1 through 6, Cafeteria Room, West Restrooms	None Detected
B-25	12" x 12" black floor tile Hallway 2, Offices 1 through 6, Cafeteria Room, West Restrooms	None Detected
B-25 Mastic	Yellow mastic Hallway 2, Offices 1 through 6, Cafeteria Room, West Restrooms	None Detected
B-26	2' x 4' ceiling tile Hallway 1, Cafeteria Room	None Detected

TABLE 1 SUMMARY OF SUSPECTED ACBM BULK SAMPLES Office Building, 9003 North Main Street, Houston, Texas

SAMPLE NO.	DESCRIPTION/LOCATION	RESULT
B-27	2' x 4' ceiling tile Hallway 1, Cafeteria Room	None Detected
B-28	2' x 4' ceiling tile Hallway 1, Cafeteria Room	None Detected
B-29	Wallboard Cafeteria Room	None Detected
B-29 JC	Joint compound/wall texture Cafeteria Room	None Detected
B-30	Wallboard Cafeteria Room	None Detected
B-30 JC	Joint compound/wall texture Cafeteria Room	None Detected
B-31	Wallboard Cafeteria Room	None Detected
B-31 JC	Joint compound/wall texture Cafeteria Room	None Detected
B-32	Interior Window Glazing Offices 6, 7, 8 and Restroom A	3 – 5% Chrysotile
B-33	Interior Window Glazing Offices 6, 7, 8 and Restroom A	Not Analyzed
B-34	Interior Window Glazing Offices 6, 7, 8 and Restroom A	Not Analyzed

Notes:

- 1) Bulk samples were analyzed by ATC's Laboratory in Miami, Florida utilizing the Environmental Protection Agency's Interim Method for the Detection of Asbestos in Bulk Insulation Samples. (EPA 600/M4-82020. July 1993) and the McCrone Research Institute's The Asbestos Particle Atlas as method references.
- 2) ATC's laboratory is accredited by the NVLAP, participates in the NVLAP and AIHA Bulk Asbestos Sample Quality Assurance Programs, and is licensed to analyze bulk asbestos samples by the Texas Department of Health.
- 3) Due to the sometimes difficulty in analyzing non-friable or resinously bound materials, ATC's laboratory recommends that these materials, which were not found to contain asbestos, be analyzed using alternative methods of identification, such as Transmission Electron Microscopy.

TABLE 2 SUMMARY OF ACBM ASSESSMENT Houston, Texas **QUANTITY** HAZARD **MATERIAL/LOCATION ESTIMATE** FRIABILITY ASSESSMENT Wallboard and Joint Compound 25,000 SF Non-friable Cove Base Mastic 2500 LF Non-friable A Carpet Mastic 650 SF Non-friable Α 1' x 1' Ceiling Tile and Mastic 775 SF Non-friable Α 12" x 12"White w/Tan Splotches floor tile and 250 SF Non-friable Α mastic 2' x 2' Ceiling Tile 350 SF Friable Α 12" x 12" Black floor tile and mastic 2,700 SF Non-friable Α 2' x 4' Ceiling Tile 132 SF Friable Α Texture, Wallboard and Joint Compound Friable 950 SF Α

232 LF

Non-friable

Notes:

- C-1 Asbestos Present Serious health hazard as defined by EPA. Abatement should be top priority.
- C-2 Asbestos Present Health hazard as defined by EPA. Abatement should be planned.
- C-3 Asbestos Present No action necessary unless renovation, remodeling, or demolition is planned.
- **B-1** Asbestos Present Contains 1% asbestos or less. Not regulated by TDSHS.
- **B-2** Asbestos Present Adequately enclosed.
- B-3 Asbestos Present Adequately encapsulated
- A No Asbestos Found

Interior Window Glazing

Offices 6, 7, 8 and Restroom A

A-1 Asbestos Abated – Once identified Asbestos-Containing Materials have been abated.

Unsampled asbestos-containing construction materials may be located within walls, ceiling cavities, below flooring or grade, and other non-accessible areas. Precaution should be used in relation to these unsampled materials until their asbestos content has been determined by proper sampling and analysis.

C-2

TABLE 3 SUMMARY OF ANALYSIS OF PAINT FOR LEAD DETERMINATION 9003 North Main Street, Houston, Texas

	The state of the s		2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
Sample No.	Sample Site	Tested Component	Analytical Results
L-001	CWB	Off-White paint on concrete block wall	< 0.02
L-002	CWB	Off-White paint on plywood	< 0.02
L-003	WHB	Off-White paint on interior plywood walls	0.02
L-004	OFB	Red Primer on structural steel	0.66
L-005	OFB	Off-White over Tan on walls at north end of warehouse	< 0.02
L-006	OFB	White paint on walls in Cafeteria Room	< 0.01
L-007	OFB	Tan paint on all office walls	< 0.01
L-008	OFB	Tan paint on all office walls	< 0.01

Notes:

CWB- Car Wash Building

WHB - Warehouse Building

OFB - Office Building

Paint chip samples were analyzed by ATC's Laboratory in New York, New York utilizing the Environmental Protection Agency's Analytical Method 7420 and Digestion Method 3050.

²⁾ ATC's laboratory is accredited by the AIHA and certified under ELLAP.

TABLE 4 SUMMARY OF LBPM ASSESSMENT 9003 North Main Street, Houston, Texas

LOCATION/DESCRIPTION	QUANTITY ESTIMATE	CONDITION	HAZARD ASSESSMENT
Off-White paint on concrete block wall	512 SF	Fair	A
Off-White paint on plywood	1280 SF	Fair	A
Off-White paint on interior plywood walls	5000 SF	Good	A
Red Primer on structural steel	2000 SF	Fair	C-1
Off-White over Tan on walls at north end of warehouse	3200 SF	Good	A
White paint on walls in Cafeteria Room	950 SF	Good	A
Tan paint on all office walls	25,000 SF	Good	A
Tan paint on all office walls	25,000 SF	Good	A

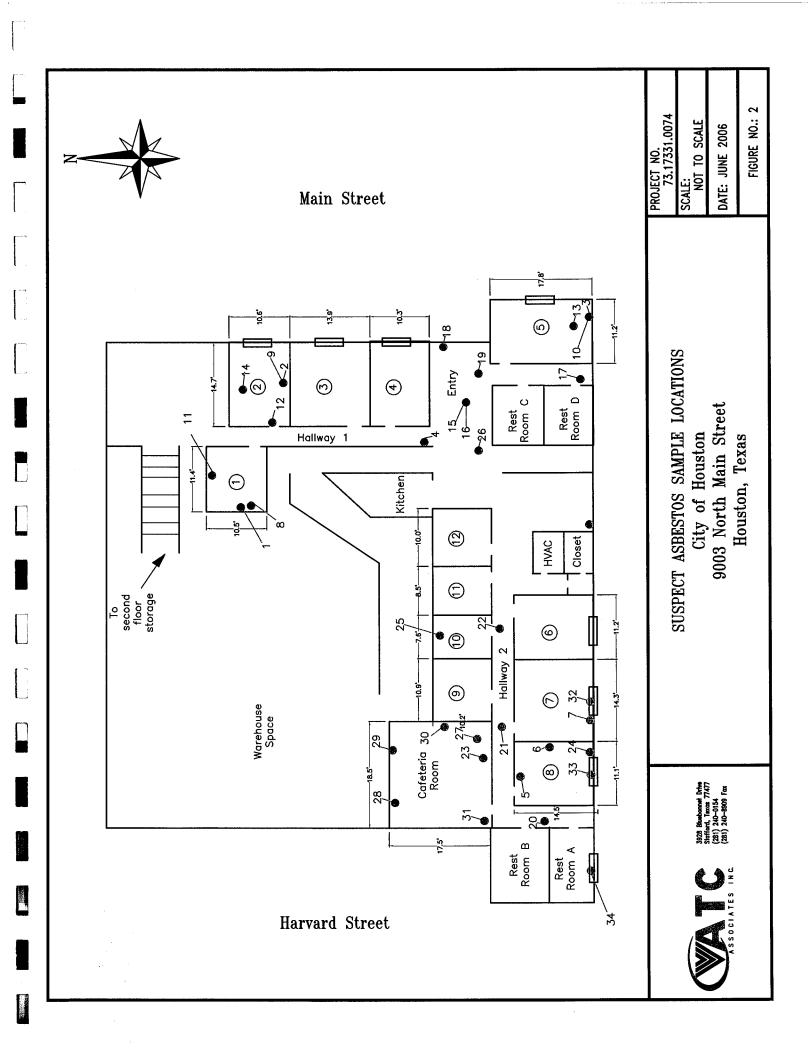
Notes:

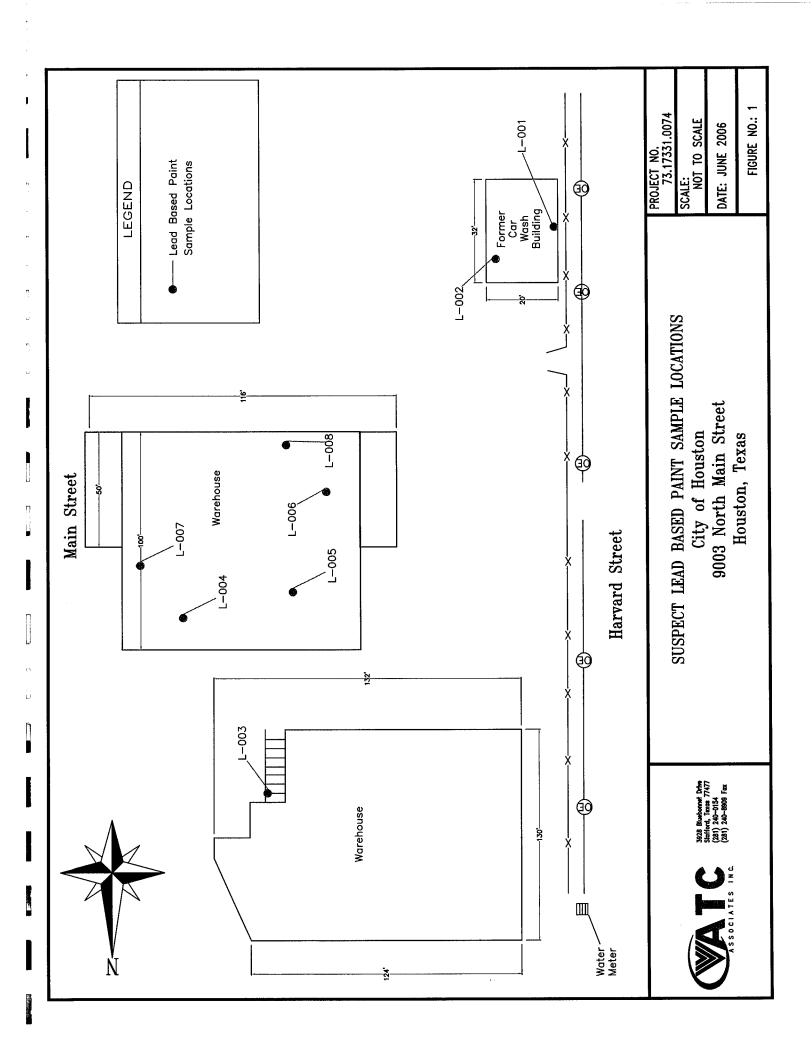
- C-1 Lead Present Health hazard as defined by applicable Federal and State regulations. Abatement priority. (≥5,000 ppm or 0.5% by weight or 1 mg/cm²)
- C-2 Lead Present No action necessary when lead levels are below applicable Federal and State regulation action levels. OSHA regulations may apply to workers during demolition or renovation. (<5,000 ppm or 0.5% by weight or 1 mg/cm²)</p>
- A Allowable Lead Level (≤600 ppm or 0.06% by weight as defined by the U.S. Consumer Product Safety Commission (CPSC) report dated October 1, 1996)
- A-1 Lead Abated Identified Lead Containing Materials (LCM) have been abated.

Un-sampled LBPM may be located within walls, ceiling cavities, below flooring or grade, and other non-accessible areas. Precaution should be used in relation to these unsampled materials until their lead content has been determined by proper sampling and analysis.



APPENDIX A SAMPLE LOCATION FIGURES







APPENDIX B

SUSPECTED ACBM BULK SAMPLE ANALYSIS REPORT AND CHAIN-OF-CUSTODY



NVLAP Accredited #200335-0

9955 NW 116th Way, Suite 1, Miami, Florida, 33178, (305)882-8200, (305)882-1200 (fax)

BULK SAMPLE ANALYSIS REPORT

Page 1 of 5

Client Name:

ATC - Houston

73.17331.0074

Task #

Batch

06-1174

Project #

City of Houston, 9003 Main Street

Date Received

6/29/2006

Project Name

Date Analyzed Analyst

6/29/2006

Date Collected

6/28/2006

Domingo Ramos

CollectedBy:

Jennifer Boone

Analyst Signature

On 06/29/06, Thirty Four(34) bulk samples were submitted by Jennifer Boone for Polarized Light Microscopy (PLM)/Dispersion Staining Analysis.

Of the Thirty Four(34) samples submitted, Fifty One(51) analyses were performed with associated layers for asbestos content utilizing EPA Method 600/R-93/116. Copies of the chain of custody data sheets are attached; additional information may be found therein. The results are summarized below.

Sample ID/	Sample	Stereosco	pic	Asbestos	% Fibrous	%	Non-Fibrous
Lab ID	Description	Description	on .	Type(s)	Compone	ents	Components
B-01	Wallboard	Color: Friable: Homo: Layered:	White, Tan Yes Yes Yes	None Detected	Cellulose Fibrous Gla	5-10 ass 2-3	Silicates
B-01	Joint Compound	Color: Friable: Homo: Layered:	White Yes Yes No	None Detected	None Dete	cted	Silicates Carbonates
B-02	Wallboard	Color: Friable: Homo: Layered:	White, Tan Yes Yes Yes	None Detected	Cellulose Fibrous Gla	5-10 ass 2-3	Silicates
B-02	Joint Compound	Color: Friable: Homo: Layered:	White Yes Yes No	None Detected	None Dete	cted	Silicates Carbonates
B-03	Wallboard	Color: Friable: Homo: Layered:	White, Tan Yes Yes Yes	None Detected	Cellulose Fibrous Gla	5-10 ass 2-3	Silicates
В-03	Joint Compound	Color: Friable: Homo: Layered:	White Yes Yes No	None Detected	None Dete	cted	Silicates Carbonates
B-04	Wallboard	Color: Friable: Homo: Layered:	White, Tan Yes Yes Yes	None Detected	Cellulose Fibrous Gla	5-10 ass 2-3	Silicates
B-04	Joint Compound	Color: Friable: Homo: Layered:	White Yes Yes No	None Detected	None Deter	cted	Silicates Carbonates
B-05	Wallboard	Color: Friable: Homo: Layered:	White, Tan Yes Yes Yes	None Detected	Cellulose Fibrous Gla	5-10 ass 2-3	Silicates
B-05	Joint Compound	Color: Friable: Homo: Layered:	White Yes Yes No	None Detected	None Detec	cted	Silicates Carbonates
B-06	Wallboard	Color: Friable: Homo: Layered:	White, Tan Yes Yes Yes	None Detected	Cellulose Fibrous Gla	5-10 ass 2-3	Silicates
B-06	Joint Compound	Color: Friable: Homo: Layered:	White Yes Yes No	None Detected	None Detec	cted	Silicates Carbonates



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BULK SAMPLE ANALYSIS REPORT

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Sample ID/	Sample	Stereoscopic	Asbestos %	Fibrous %	Non-Fibrous
Lab ID	Description	Description	Type(s)	Components	Components
B-07	Wallboard	Color: White, Tan Friable: Yes Homo: Yes Layered: Yes	None Detected	Cellulose 5-10 Fibrous Glass 2-3	Silicates
B-07	Joint Compound	Color: White Friable: Yes Homo: Yes Layered: No	None Detected	None Detected	Silicates Carbonates
B-08	Covebase Mastic	Color: Tan Friable: No Homo: Yes Layered: No	None Detected	None Detected	Silicates Mastic
B-09	Covebase Mastic	Color: Tan Friable: No Homo: Yes Layered: No	None Detected	None Detected	Silicates Mastic
B-10	Covebase Mastic	Color: Tan Friable: No Homo: Yes Layered: No	None Detected	None Detected	Silicates Mastic
B-11	Carpet Mastic	Color: Yellow Friable: No Homo: Yes Layered: No	None Detected	None Detected	Silicates Mastic
B-12	Carpet Mastic	Color: Yellow Friable: No Homo: Yes Layered: No	None Detected .	None Detected	Silicates Mastic
B-13	Carpet Mastic	Color: Yellow Friable: No Homo: Yes Layered: No	None Detected	None Detected	Silicates Mastic
B-14	1' x 1' Ceiling Tile	Color: White Friable: Yes Homo: Yes Layered: No	None Detected	Cellulose 1-2 Fibrous Glass 65-70	Silicates Perlite NF Glass
B-14	Brown Mastic	Color: Brown Friable: No Homo: Yes Layered: No	None Detected	Fibrous Glass 3-5	Silicates Mastic
B-15	1' x 1' Ceiling Tile	Color: White Friable: Yes Homo: Yes Layered: No	None Detected	Cellulose 1-2 Fibrous Glass 65-70	Silicates Perlite NF Glass
B-15	Brown Mastic	Color: Brown Friable: No Homo: Yes Layered: No	None Detected	Fibrous Glass 3-5	Silicates Mastic



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On 06/29/06, Thirty Four(34) bulk samples were submitted by Jennifer Boone for Polarized Light Microscopy (PLM)/Dispersion Staining Analysis.

Sample ID/	Sample	Stereoscopic	Asbestos	%	Fibrous	%	Non-Fibrous
Lab ID	Description	Description	Type(s)		Components		Components
B-16	1' x 1' Ceiling Tile	Color: White Friable: Yes Homo: Yes Layered: No	None Detected		Cellulose Fibrous Glass	1-2 65-70	Silicates Perlite NF Glass
B-16	Brown Mastic	Color: Brown Friable: No Homo: Yes Layered: No	None Detected		Fibrous Glass	3-5	Silicates Mastic
B-17	12 x 12 White Floor Tile w/ Tan Splotches	Color: White, Friable: No Homo: Yes Layered: No	Yan None Detected		None Detected		Silicates Carbonates
B-17	Mastic	Color: Yellow Friable: No Homo: Yes Layered: No	None Detected		None Detected		Silicates Mastic
B-18	12 x 12 White Floor Tile w/ Tan Splotches	Color: White, Friable: No Homo: Yes Layered: No	Fan None Detected		None Detected		Silicates Carbonates
B-18	Mastic	Color: Yellow Friable: No Homo: Yes Layered: No	None Detected		None Detected		Silicates Mastic
B-19	12 x 12 White Floor Tile w/ Tan Splotches	Color: White, Friable: No Homo: Yes Layered: No	an None Detected		None Detected		Silicates Carbonates
B-19	Mastic	Color: Yellow Friable: No Homo: Yes Layered: No	None Detected		None Detected		Silicates Mastic
B-20	2' x 2' Ceiling Tìle	Color: Brown, Friable: Yes Homo: Yes Layered: No	White None Detected		Cellulose Fibrous Glass	20-25 10-15	Silicates Perlite NF Glass
B-21	2' x 2' Ceiling Tile	Color: Brown, Friable: Yes Homo: Yes Layered: No	White None Detected		Cellulose Fibrous Glass	20-25 10-15	Silicates Perlite NF Glass
B-22	2' x 2' Ceiling Tile	Color: Brown, Yes Homo: Yes Layered: No	White None Detected		Cellulose Fibrous Glass	20-25 10-15	Silicates Perlite NF Glass
B-23	12 x 12 Black Floor Tile	Color: Black Friable: No Homo: Yes Layered: No	None Detected		None Detected		Silicates Carbonates

NVLAP Accredited #200335-0

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BULK SAMPLE ANALYSIS REPORT

Page 4 of 5

Client Name:

ATC - Houston

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Sample ID/	Sample	Stereoscopic	Asbestos %	Fibrous %	Non-Fibrous
Lab ID	Description	Description	Type(s)	Components	Components
B-23	Mastic	Color: Tan Friable: No Homo: Yes Layered: No	None Detected	None Detected	Silicates Mastic
B-24	12 x 12 Black Floor Tile	Color: Black Friable: No Homo: Yes Layered: No	None Detected	None Detected	Silicates Carbonates
B-24	Mastic	Color: Tan Friable: No Homo: Yes Layered: No	None Detected	None Detected	Silicates Mastic
B-25	12 x 12 Black Floor Tile	Color: Black Friable: No Homo: Yes Layered: No	None Detected	None Detected	Silicates Carbonates
B-25	Mastic .	Color: Tan Friable: No Homo: Yes Layered: No	None Detected	None Detected	Silicates Mastic
B-26	2' x 4' Ceiling Tile	Color: White, Tan Friable: Yes Homo: Yes Layered: No	None Detected	None Detected	Silicates Perlite
B-27	2' x 4' Ceiling Tile	Color: White, Tan Friable: Yes Homo: Yes Layered: No	None Detected	None Detected	Silicates Perlite
B-28	2' x 4' Ceiling Tile	Color: White, Tan Friable: Yes Homo: Yes Layered: No	None Detected	None Detected	Silicates Perlite
B-29	Wallboard	Color: White, Tan Friable: Yes Homo: Yes Layered: Yes	None Detected	Cellulose 5-10 Fibrous Glass 2-3	Silicates
B-29	Texture/Joint Compound	Color: White Friable: Yes Homo: Yes Layered: Yes	None Detected	None Detected	Silicates Carbonates Paint
B-30	Wallboard	Color: White, Tan Friable: Yes Homo: Yes Layered: Yes	None Detected	Cellulose 5-10 Fibrous Glass 2-3	Silicates
B-30	Texture/Joint Compound	Color: White Friable: Yes Homo: Yes Layered: Yes	None Detected	None Detected	Silicates Carbonates Paint



NVLAP Accredited #200335-0

9955 NW 116th Way, Suite 1, Miami, Florida, 33178, (305)882-8200, (305)882-1200 (fax)

BULK SAMPLE ANALYSIS REPORT

Page 5 of 5

Client Name:

ATC - Houston

Project #
Project Name

73.17331.0074

City of Houston, 9003 Main Street

Task #

Batch

06-1174

Date Received

6/29/2006

Date Analyzed

6/29/2006

Are almos

----- D-----

Date Collected

6/28/2006

Analyst

Domingo Ramos

CollectedBy:

Jennifer Boone

Analyst Signature

Deser 5

On 06/29/06, Thirty Four(34) bulk samples were submitted by Jennifer Boone for Polarized Light Microscopy (PLM)/Dispersion Staining Analysis.

Of the Thirty Four(34) samples submitted, Fifty One(51) analyses were performed with associated layers for asbestos content utilizing EPA Method 600/R-93/116. Copies of the chain of custody data sheets are attached; additional information may be found therein. The results are summarized below.

Sample ID/ Lab ID	Sample Description	Stereoscopic Description	Asbestos Type(s)	%	Fibrous Components	%	Non-Fibrous Components
B-31	Wallboard	Color: White Friable: Yes Homo: Yes Layered: Yes	, Tan None Detected			5-10 2-3	Silicates
B-31	Texture/Joint Compound	Color: White Friable: Yes Homo: Yes Layered: Yes	None Detected		None Detected		Silicates Carbonates Paint
B-32	Interior Window Glazing	Color: Gray Friable: Yes Homo: Yes Layered: No	Chrysotile	3-5	None Detected		Silicates Carbonates
B-33	Interior Window Glazing	Color: Friable: Homo: Layered:	*Not Analyzed				
B-34	Interior Window Glazing	Color: Friable: Homo: Layered:	*Not Analyzed				·

Comments- Analytical Methods: EPA 600/R-93/116, July 1993

- Enclosed test results relates only to items tested.
- This report shall not be reproduced, except in full without written approval of the laboratory.
- This report cannot be used to claim product endorsement by NVLAP or any agency of the U.S. Government.

-Method Limitations:

Analysis of resinous and bituminous bound materials (i.e. floor tile, roofing, etc.) by Polarized Light Microscopy (PLM) may yield false negative results due to method limitations. In these cases, EPA and Hygeia recommend alternative methods of analysis.

- Layered Samples:

Samples that contain discreetly identifiable layers will be analyzed and reported separately, if any layer is found to contain asbestos. When all layers are found not to contain asbestos one composite analysis can be reported. In addition, samples that contain individual layers that cannot be discreetly separated without compromising some layer; these samples will be analyzed as composite.

-Sample Archival:

Hygeia's policy is to dispose of all unused portions of the sample(s) 90 days after analysis. Samples can be returned to the client if prior arrangements are made.

*Not Analyzed - 1st Positive Stop Series

Respectfully Submitted,

Julio Lopez

Laboratory Manager

ASBESTOS BULK SAMPLING - CHAIN OF CUSTODY

ATC ASSOCIATES INC.
3928 Bluebonnet Drive
Stafford, Texas 77477
(281) 240-0154 phone (281) 240-8909 fax

Project No.: 73.17331.0074

City of Houston

Client Name:

Date:

6/28/2006

Site Location:	ation:	9003 Main Street	Prj Manage	Manager: Catherine McLain	Inspector:	J. Boone
Sample No.	HA No.	Material Description	Classification	Homogeneous Material Location(s)	Quantity	Condition
10-07 10-04 10-04	\$	Wallboard and Jaint Compound	Surfacing TSI Miscellaneous Nonfriable Friable	Office 1-west wall Office 2-South wall Office 5-South wall	25,000 SF	Good Fair Poor
Doct Book			Surfacing TSI Miscellaneous Nonfriable Friable	the South End of the Chief of the Shice of North wall		Good Fair Poor
200	>		Surfacing TSI Miscellaneous Nonfriable Friable	Stice 8 - East Wall Stice 7 - Suth Wall		Good Fair Poor
B-08 B-09	(\mathcal{A})	Cove Base Mastic	Surfacing TSI Miscellaneous Nonfriable Friable	Office 1 - West Well Office 2 - South Well Office 5 - South Well	03500 6 F	Good Fair Poor
12 PM	0	Carpet Mastic	Surfacing TSI Miscellaneous Nontriable Friable	Office 1 - North wall Office 2 - Dooth wall-doorway	15059	Good Fair Poor
1000 1000 1000	\bigcirc	1'x1'Cerling The and Frown Wastic	Surfacing TSI Miscellaneous Nonfriable Friable	Office 2 - Center	78 STM	Good Fair Poor
B-187	W	12x12 white wo Taxion of Janie 2015 takes - foothie	Surfacing TSI Wiscellaneous Nonfriable Friable	Entry Hall Buthroom Dovway Entry Hall-Front Rorway Caty tall-tall Dorway	350 SF	Good . Fair Poor
COMMENTS:	TS:	Positive stop analysis per homogenous sample.	genous sample.			
RELINQUI	SHED BY	RELINQUISHED BY (SIGNATURE BELOW):	RECEIVED BY (S	RECEIVED BY (SIGNATURE BELOW):	LABORATORY:	TurnAround
PRINT'NAME:	WE:	DATE/TIME:	PRINT NAME:	DATE/TIME:	·	48 hours 24 Hours Other
Jenn	Jenniterh	Doone 6/28/06/0630pm	Om Cago	05:01 70/62/9		

ASBESTOS BULK SAMPLING - CHAIN OF CUSTODY

ATC ASSOCIATES INC. 3928 Bluebounet Drive Stafford, Texas 77477 (281) 240-0154 phone (281) 240-8909 fax Project No.: 73.17331.0074

City of Houston

Client Name:

Site Location: 9003 Main Street

Prj Manager: Catherine McLain

6/28/2006 Date:

Inspector: J. Boome

Sample No.	HA No.	Material Description	Classification	Homogeneous Material Location(s)	Quantity	Condition
877 877		21×21 Ceoling Tule	Surfacing TSI Miscellaneous Nontriable Friable	Hallway 2 Fallway 2 Fallbay 3	350 SF	Good Fair X Poor
0,000 0,000 0,000		12x12 Black footh	Surfacing 7SI Miscellaneous Nonfriable Friable	Captair Com	2700 SF	Good Fair Poor
0-26 0-27 0-27 0-23	#	24 Gelling File	Surfacing TSI Miscellaneous Nonfriable Friable	Sallway 1- south end Capetiticalor . West without Capetina Bon - North end	1325F	Good Fair X Poor
B-29 B-30 B-31	H	Tecture/Ub/11 board	Surfacing 7SI Miscellaneous Nonfriable Friable	Cathria Room - Northibel East-Uest	950SF	Good CX Fair Poor
82.00 52.00 52.00 52.00	1	Windows glazing-	Surfacing 7SI Miscellaneous Nonfriable Friable	Opin of Opins of Section A.	232 LF	Good Fair Poor
)			Surfacing 7SI Miscellaneous Nonfriable Friable			Good Fair Poor
			Surfacing 7SI Miscellaneous Nonfriable Friable			Good Fair Poor
COMMENTS:	S:	Positive stop analysis per homogenous	genous sample.			



Standard
48 hours
24 Hours
Other

DATE/TIME:

PRINT NAME:

DATE/TIME: Dale Jane

PRINT NAME:

10630911

TurnAround

LABORATORY:

RECEIVED BY (SIGNATURE BELOW)

RELINQUISHED BY (SIGNATURE BELOW):



APPENDIX C

SUSPECTED LEAD-BASED PAINT ANALYSIS REPORT AND CHAIN-OF-CUSTODY



ATC ASSOCIATES INC

104 E. 25th Street, 10th Floor New York, NY 10010 Tel. 212-353-8280 Fax: 212-353-8306

REPORT DATE:

6/30/2006

CLIENT NAME:

ATC - TEXAS

PROJECT NAME:

CITY OF HOUSTON / 9003 MAIN

SAMPLED BY:

Client

SAMPLE DATE:

6/28/2006

RECEIVED DATE:

6/30/2006

ANALYZED DATE:

6/30/2006

SAMPLE MEDIA:

Paint Chips by %

ANALYSIS REQUIRED:

Method EPA 3050/7420

SAMPLE ID	BATCH NO.		LEAD CONCENTRATION (% by weight)	DETECTION LIMIT (% by weight)
. L-001	-	39613	<0.02	0.02
L-002		39613	<0.02	0.02
L-003		39613	0.02	0.02
L-004		39613	0.66	0.01
L-005		39613	<0.02	0.02
L-006		39613	<0.01	0.01
L-007		39613	<0.01	0.01
L-008		39613	<0.01	0.01

SAMPLE ID

BATCH NO.

LEAD CONCENTRATION (% by weight)

DETECTION LIMIT (% by weight)

NOTE 1:

THE REPORTING LIMIT (RL) IS 0.01%. The detection limit as reported is the reporting limit. The true detection limit is half the RL.

HUD defines lead-based paint as paint having a lead concentration equal to or above 0.5% by weight.

Results preceded by "<" are below the detectable levels by this analysis method.

ATC Associates Inc. Laboratory is not responsible for sample collection. These results relate only to the items tested. This report shall not be reproduced, except in full, without written approval of the laboratory.

CERTIFICATION	NS AND	SIGNA'	TURES
---------------	--------	--------	-------

ANALYSIS PERFORMED BY:

Inna Livshiz

REPORT PREPARED BY:

Inna Kipen

ATC certifies that this report is an accurate and authentic report of results obtained from the laboratory analysis.

QUALITY ASSURANCE COORDINATOR: Mei Wang

LABORATORY DIRECTOR:

Milena Lowd

ATC is accredited by the New York State Department of Health Environmental Laboratory Approval Program (ELAP) and by the American Industrial Hygiene Association (AIHA) to perform analysis of lead in paint, dust wipes, air and soil samples. (ELAP #10879), (AIHA #100229).

The document(s) contained herein are confidential and privileged information, intended for the exclusive use of the individual or entity named above.

Unless otherwise indicated, no blank corrections were performed.

The condition of all samples was acceptable upon receipt.

Unless otherwise indicated all QC results were in control.

APPENDIX D ATC'S APPLICABLE LICENSES AND CERTIFICATIONS



Seller.

TEXAS DEPARTMENT OF STATE HEALTH SERVICES

Be it known that

ATC ASSOCIATES, INC.

is certified to perform as a

Asbestos Consultant Agency

so long as this license is not suspended or revoked and is renewed according to the in the State of Texas within the purview of Texas Occupations Code, chapter 1954, rules adopted by the Texas Board of Health.

STORY NO

Eduardo J. Sanchez, M.D., M.P.H Commissioner of Health

> License Number. 100032 Effective Date: 2/15/2006

Expiration Date: 2/14/2008

(Void After Expiration Date)



TEXAS DEPARTMENT OF STATE HEALTH SERVICES

Be it known that

ATC GROUP SERVICES DBA HYGEIA LABORATORIES

is certified to perform as a

Asbestos Laboratory TEM, PLM, PCM

so long as this license is not suspended or revoked and is renewed according to the in the State of Texas within the purview of Texas Occupations Code, chapter 1954, rules adopted by the Texas Board of Health.

SUSILINY MO

Eduardo J. Sanchez, M.D., M.P.H Commissioner of Health

> License Number 300230 Effective Oate: 5/22/2006

Expiration Date: 5/21/2008

(Void After Expiration Date)

TEXAS Department of State Health Services

Control No: 93012

Department of State Health Services certifies that:

CATHERINE G MCLAIN

is Licensed as an: Individual Asbertos Consultant

License Number: 105451 From: 05/19/2006 To: 05/18/2008



Control No:* * 87205

Department of State Health Services certifies that: JENNIFER L BOONE

is Licensed as an:

Individual Asbestos Consultant

License Number: 105554

From: To:

D

06/29/2005 06/28/2007

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Appendix B Laboratory Reports and Chain-of-Custody Documentation

EXECUTIVE SUMMARY

ATC Associates completed a limited Phase II Environmental Site Assessment (ESA) of the City of Houston property located at 9003 North Main Street, in Houston, Texas (herein referred to as the subject property). This report details the completion of the Limited Phase II ESA, in accordance with applicable ASTM standards and accepted environmental practices.

This Limited Phase II ESA conducted on June 21, 2006 identified the presence of BTEX and formaldehyde in the soil and groundwater. The constituent concentrations identified during the Phase II ESA were evaluated with respect to previous concentrations associated with the leaking petroleum storage tank (LPST) case and the TRRP Tier 1 groundwater protective concentration levels (PCLs). BTEX concentrations detected during this assessment were above TCEQ action levels for petroleum underground storage tank sites. These concentrations were below the previous concentrations detected at the site when closure was granted in 1997. The evaluation also identified formaldehyde to be below the TRRP Tier 1 groundwater PCL.

Three soil borings were advanced during the course of this Limited Phase II ESA. Soil samples were collected at the 12 – 14 ft below ground surface (bgs) interval in B-1, and the 20-22 ft bgs interval in B-2 and B-3. One soil samples from each boring was analyzed for BTEX by EPA Method 8021B, TPH by TX Method 1005, and Formaldehyde by EPA Method 8315. Analytical results for BTEX and TPH were below previous concentrations under which the site closed and formaldehyde was below the TRRP Tier 1 Residential PCLs.

The soil borings were converted to temporary monitoring wells. Groundwater samples were collected, submitted to Severn Trent Laboratories (STL) and analyzed for BTEX by EPA Method 8021B, TPH by TX Method 1005, and Formaldehyde by EPA Method 8315.

The concentrations identified during the Phase II ESA were evaluated with respect to previous concentrations associated with LPST case and the TRRP Tier 1 groundwater protective concentration levels (PCLs). BTEX was detected above TCEQ action levels but were below previous concentrations at the site. Formaldehyde concentrations were below the TRRP Tier 1 Residential PCLs.

Based on the data provided during the course of this Limited Phase II ESA, it appears that no further actions are recommended for the subject property at this time.

1.0 INTRODUCTION

1.1 Purpose and Scope

ATC Associates (ATC) was contracted by the City of Houston to conduct a Limited Phase II ESA of the subject property located at 9003 North Main in Houston, TX.

The purpose of the Limited Phase II Environmental Site Assessment (ESA) is to assess impact to the property from the LPST release associated with the site as well as reported historical use as a funeral home. The Limited Phase II ESA was performed to assist in confirming the nature and extent of contamination in the soil and groundwater of the above-referenced property.

The scope of the Limited Phase II Investigation included subsurface soil sampling, the installation of three temporary groundwater monitoring points, and groundwater sampling via the installation of three soil borings and temporary monitoring wells.

1.2 Background

The property is owned by the City of Houston, and was formerly used as an office and storage facility by the Street Maintenance Department. The site contains a one-story office building with an attached warehouse area in the eastern section of the property, a metal warehouse building at the north end of the property, and a canopy covered truck wash area on the southwestern portion of the site. The site also contains a concrete paved storage area formerly used for gravel, soil and other bulk materials used for street maintenance. The subject property was also identified in the TCEQ database as a leaking petroleum storage tank (LPST) site. A total of four USTs located at the property were reportedly removed in 1992.

2.0 PREVIOUS INVESTIGATIONS

2.1 Phase I Environmental Site Assessments

Weston conducted a Phase I ESA which identified the following environmental concerns:

- The subject property was identified in the TCEQ database as a leaking petroleum storage tank (LPST) site. A total of four USTs located at the property were reportedly removed in 1992. Following removal of the USTs, an assessment and groundwater monitoring activities were performed at the site. Groundwater was impacted but it was determined that there were no apparent threats or impacts to receptors. TCEQ issued final closure of the LPST case in February 1998.
- The office building present at the property was constructed prior to the 1970s and may contain asbestos and lead-based paint.
- The property was reportedly used as a funeral home prior to 1984; however, no documentation is available to confirm the former presence of a funeral home on the site.

2.2 Phase II Environmental Site Assessment

Carter and Burgess conducted Site Assessment activities at the site under the TCEQs LPST program in June 1995. Three groundwater monitoring wells were installed and soil and groundwater sampling was conducted. The soil and groundwater concentrations were above TCEQ action levels and the site was issued LPST number 104846. Groundwater monitoring was conducted and the site was closed in December 1997.

3.0 SOIL ASSESSMENT

3.1 Soil Boring Advancement and Sample Collection

On June 21, 2006, three soil borings were advanced with push-probe drilling equipment in the areas of the former USTs and near the back door of the office building. A site map depicting the soil boring location is included as Figure 1. The soil boring logs are provided in Appendix B.

During the advancement of the soil borings, soils were sampled continuously every 2 feet. One soil sample from each soil boring was selected for analysis; either the sample with the highest OVM reading or the sample at the soil groundwater interface was collected for laboratory analysis. Each soil sample was analyzed for benzene, toluene, ethylbenzene, and xylenes (BTEX) by EPA Method 8021B, total petroleum hydrocarbons (TPH) by Texas Method 1005, and formaldehyde by EPA Method 8315.

3.2 Laboratory Analysis

Analytical results of the soil samples collected by ATC indicated the presence of BTEX, and formaldehyde. Benzene concentrations ranged from below laboratory method detection limits in boring B-1 and B-3 to 0.072 mg/kg in B-1. Total BTEX concentrations ranged from below laboratory method detection limits in borings B-1 and B-3 to 2.982 mg/kg in B-2. TPH results were below laboratory method detection limits in the samples that were submitted for analysis. Formaldehyde was detected at concentrations of 2.520 mg/kg in B-1, 0.575 mg/kg in B-2, and 0.357 mg/kg in B-3. A copy of the laboratory analytical report by STL is provided in Appendix B.

The benzene concentration detected in B-2 is above the TCEQ action level for a UST site but is below the concentrations under which the site closed in 1997. The formaldehyde concentrations are below the TRRP Tier 1 Residential PCLS.

4.0 GROUNDWATER ASSESSMENT

4.1 Groundwater Sampling

The soil borings were converted to temporary groundwater monitoring wells. The temporary monitoring wells were constructed of threaded connection 1-inch ID, Schedule 40 PVC solid pipe, and 0.010-inch slotted PVC well screen.

On June 21, 2006, the three temporary monitoring wells were purged and sampled using a peristaltic pump. The wells were purged prior to sampling to reduce the amount of sediment present in the groundwater samples.

4.2 Laboratory Analysis

The analytical results of the groundwater samples obtained by ATC from the temporary monitoring wells on June 21, 2006 indicated concentrations of BTEX and formaldehyde. Benzene concentrations ranged from below laboratory method detection limits in B-2 and B-3 to 25.6 ug/L in B-1. Total BTEX concentrations ranged from below laboratory detection limits in B-3 to 561 ug/L in B-1. Formaldehyde concentrations were 27.6 ug/L in B-1, 46.4 ug/L in B-2, and 9.8 ug/L in B-3. TPH concentrations were below laboratory detection limits in the three borings. A copy of the laboratory analytical report is included in Appendix B

5.0 REGULATORY EVALUATION

Analytical laboratory results from this Limited Phase II ESA were compared to action levels established by the TCEQ and to applicable TRRP PCL. In addition, BTEX and TPH concentrations were compared to previous concentrations from when the site was in the TCEQ LPST program.

5.1 Soils

The benzene concentration detected in B-1 is above the TCEQ action levels for UST sites but is less than previous concentrations under which the site closed in 1997. The formaldehyde concentrations detected in the groundwater samples collected from the temporary monitoring wells was below the TRRP Tier I Residential PCLs.

5.2 Groundwater

The benzene concentration detected in B-1 is above the TCEQ action levels for UST sites but is less than previous concentrations under which the site closed in 1997. The formaldehyde concentrations detected in the groundwater samples collected from the temporary monitoring wells was below the TRRP Tier I Residential PCLs.

6.0 QUALITY CONTROL/QUALITY ASSURANCE

6.1 Decontamination Procedures

Drill operations were conducted using hydraulic direct push rig, with plastic sleeves, which were replaced after each 4-foot push. The auger drill rig used metal samplers which were cleaned in an alconox solution after each sample.

6.2 Field QA/QC Procedures

Soil samples submitted for laboratory analysis were contained within a Teflon-lined glass jar, placed on ice, and transported to the laboratory for analysis. Soil samples submitted to the laboratory were analyzed for BTEX, TPH and formaldehyde.

The groundwater samples submitted for laboratory analysis were contained within the appropriate containers, placed on ice, and transported to the laboratory for analysis. The groundwater sample submitted to the laboratory was analyzed for BTEX, TPH and formaldehyde.

Each sample was labeled and secured to preserve the integrity of the identification, from the time the sample was collected until it was opened at the laboratory. For each sample, the sample container label and chain-of-custody form were completed. Soil and groundwater samples were immediately placed in a cooler containing ice or frozen ice packs and hand delivered to the laboratory.

6.3 Sample Quality Control/Quality Assurance

Various QA/QC procedures were followed by the environmental laboratory. Prior to initiating analysis, it is required to establish that a given instrument meets the method tuning standard. The calibration of each instrument was verified at frequencies specified in the EPA approved methods. A new standard curve must be prepared as specified in each method per EPA Method SW-846.

Prior to analysis, instruments are required to be calibrated by the appropriate procedure. Each calibration standard was tabulated and the retention times recorded. The laboratory QA/QC results are provided in Appendix C.

7.0 CONCLUSIONS AND RECOMMENDATIONS

7.1 Conclusions

ATC has provided the following conclusions of this Phase II Site Assessment based on the field activities conducted on June 21, 2006 at the subject property, and on laboratory analytical data of media samples collected by ATC.

- The soils at the site consist of silty clays. Groundwater was encountered at a depth of approximately 15 to 21 feet bgs. Three temporary monitoring wells were installed to a maximum depth of 25 feet bgs.
- Analytical results of soil samples collected during the advancement of three soil borings along the west/southwest property boundary indicated the presence of BTEX and formaldehyde
- Formaldehyde concentrations in soil and groundwater were below the TRRP Tier I Residential PCLs

Project No. 73.27434.0032 ATC Associates Inc.

7.2 Recommendations

Based on the analytical results of soil and groundwater samples collected during this Limited Site Assessment, no further action is recommended at this time. Even though the Benzene concentrations were above TCEQ action levels, they were below the previous soil and groundwater concentrations detected at the site under which site closure was achieved.

TABLES

TABLE 1 SOIL ANALYTICAL RESULTS City of Houston 9003 North Main Houston, Texas

			Domoznod	Tologo	Cebalkonana	Joneph. A	Total BTEV	L	Fotal Petroleum Hydrocarbons	a Hydrocarbor	SL	
Well	Date	Date Depth (feet)	(mg/kg)	(mg/kg)	(mg/kg) (mg/kg) (mg/kg)	Ayienes (mg/kg)	(mg/kg)	C6-C12	i	C6-C28	Total	Formaldehyde
			(99)	/BB	(G., G.,)	ê .	(9	(mg/kg)	(mg/kg)	(mg/kg)	C6-C35	(mg/kg)
B-1	06/21/06	06/21/06 12-14	<0.0056	<0.00774	<0.00742	<0.0233	QN	<6.57	<13.8	<13.8	<13.8	2.520
B-2	B-2 06/21/06 20-22	20-22	0.0721	0.155	0.785	1.970	2.982	<6.32	<13.3	<13.3	<13.3	0.575
B-3	B-3 06/21/06	20-22	<0.00539	<0.00745	<0.00715	<0.0224	ND	<6.33	<13.3	<13.3	<13.3	0.357

Notes:

mg/kg - milligrams per kilogram or parts per million.

BTEX - benzene, toluene, ethylbenzene, and xylenes by EPA Method 8021B.

ND - not detected above laboratory detection limits.

NA - not available.

TABLE 2 GROUNDWATER ANALYTICAL RESULTS City of Houston 9003 North Main Houston, Texas

		Renzene	Toluono	T the lates	V.40	Tatal Park	L	Total Petroleum Hydrocarbons	Hydrocarbon	SI	
Well	Date	(ng/L)	(ug/L)	(ug/L)	(ug/L)	(ue/L)	C6-C12	C12-C28	C6-C28	Total	Formaldehyde
				,	, D		(mg/L)		(mg/L)	C6-C35	(ng/L)
B-1	06/21/06	25.6	7.1	131	396.9	561	<0.48	<0.83	<0.83	<0.83	27.6
B-2	90/17/90	<0.170	<0.190	0.67	2.37	3.40	<0.48	<0.83	<0.83	<0.83	46.4
B-3	06/21/06	<0.170	<0.190	<0.230	<0.560	QN	<0.48	<0.84	<0.84	<0.84	8.6

Notes:

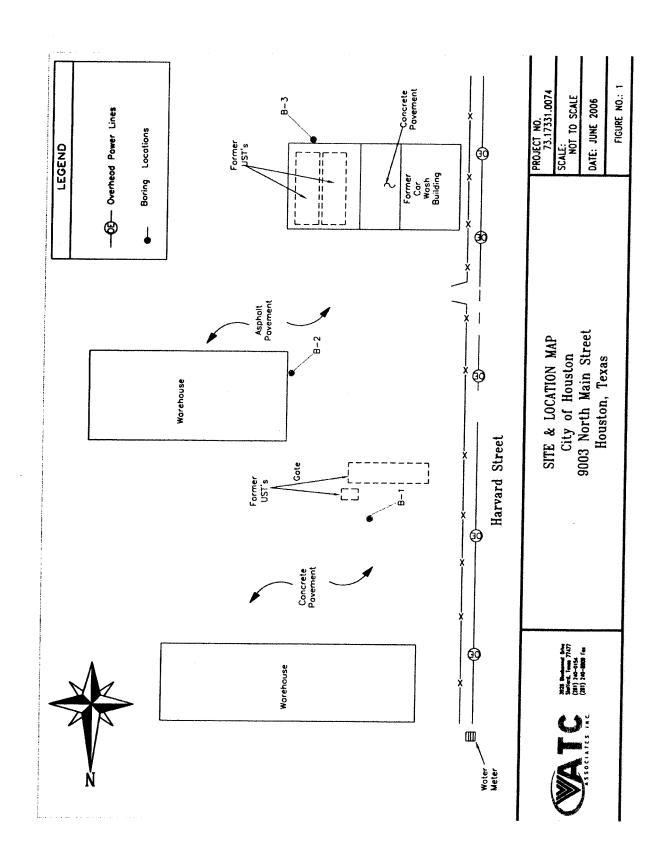
mg/kg - milligrams per kilogram or parts per million.

BTEX - benzene, toluene, ethylbenzene, and xylenes by EPA Method 8021B.

ND - not detected above laboratory detection limits.

NA - not available.

FIGURES



APPENDIX A **SOIL BORING LOGS**

	JECT► COH 9003 N. M		Bori	119							331.0074
	GED BY► PD		110	\cap	START					} 	
	CKED BY► UND SURFACE			9	COMP						
ELEV	/ATION DATUM (FT-	MSL) ►		DRIL	ING C	OMP	ANY	► Alp	ine		
DRIL	LING EQUIPMENT >			DRIL	.ER►						
BOR	ING DEPTH(FT)►	WELL DEPTH(FT) >	WATE	R DEF	PTH(F	T)-I	nitia	l:		Complete:
WELL	L MATERIALS ►			OVM,	OVA >	- OVM					
BAC	KFILL MATERIAL►										
Œ	LI	THOLOGY		***************************************	·····			S	AMPL	F	
						T	∢	1		T	
DEPTH	DE	SCRIPTION				SSGRAPHIC	OVM/OVA (PPM)	RECOVERY %		NUMBER	COMMENTS
						S. A.	Žď.	ZEC(TIME	∑ 2	
0	Concrete Gravel and Sand fill					300	3	-			
	Brown and gray silty clay										
							13				
5											
	-mixed with sand										
							2				
10	Brown and gray silty clay			**							(12–14')
							14				(,_ ,,
	Gray sand wet • 13'		T				,,,,,				
15	Brown and gray silty clay			*****			9				
	• • • •										
\exists											
, =							0				
20	End Of Boring ● 20'					u II X			L	1	·
- =	GW sample collected										
\exists											
,_=											
25											
=											
, =											
20 — J											
POI	RING DESIGNATION			Γ					F	AGE	NUMBER

PROJECT► COH 9003 N. Main		Bori	na	PROJE	CT I	NUM	BER.	- 7:	3.173	331.0074
LOGGED BY► PD			` ' ' ' ' ' ' ' ' ' ' ' ' ' ' ' ' ' ' '	START	DAT	E >	06-2	1-06)	
CHECKED BY>			9	СОМРІ	ETIC	ON D	ATE	>		
GROUND SURFACE ELEVATION DATUM (FT-MS	SL) ►		DRIL	ING C	ОМР	ANY	► Alp	ine		
DRILLING EQUIPMENT>			DRIL	ER►						
BORING DEPTH(FT)► \	WELL DEPTH(F	T) ►	WATE	R DEP	TH(F	T)-I	nitia	l:		Complete:
WELL MATERIALS ►			OVM,	OVA ►	OVM					
BACKFILL MATERIAL ►					······	,	,			
E LITH	IOLOGY				T		1	AMPL	E	-
O DE	RIPTION				GRAPHIC	OVM/OVA (PPM)	RECOVERY %	TIME	NUMBER	COMMENTS
Concrete					909	0				
_										
						•				
Dark gray, silty clay Gray, silty clay— moist few Ca						0				
Gray, silty clay— moist few Ca	Nodules									
						0				
10										
Gray and brown clay— moist, fir	rm, Fe stains									
=						0				
15										
Red and gray clay						0				
20-										(20–22')
With sand seams— wet										(== ,
=======================================						0				
End Of Boring © 24'			·	E		0				
25 GW sample collected										
_										
30-										
BORING DESIGNATION B-2		Ä.	T C		c.			F		NUMBER OF 1

		T								
PRO	JECT► COH 9003 N. Main	Bori	nal	PROJE	ECT	NUM	BER.	> 7.	3.173	31.0074
LOG	GED BY► PD	1		START	DAT	ΓE ►	06-2	1-06		
	CKED BY►	LU	<u>y</u>	СОМР	LETIC	ON D	DATE	>		
GRO ELE	UND SURFACE VATION DATUM (FT-MSL)►		DRIL	ING C	OMP	ANY	► Alp	ine		
	LING EQUIPMENT>		DRIL	_ER ►						
BOR	ING DEPTH(FT)► WELL DEPTH(F	-T) ►	WATE	R DEP	тн(г	T)-I	nitia	l:		Complete:
WEL	L MATERIALS ►		OVM,	OVA ►	OVM					
BAC	KFILL MATERIAL►									
(F	LITHOLOGY						i	AMPL	E.	
о рертн (DESCRIPTION				GRAPHIC	OVM/OVA (PPM)	RECOVERY %	TIME	NUMBER	COMMENTS
_	Asphalt Brown and gray silt				99	0				
_										
	Gray, silty clay, moist, soft									
10		TO THE TENNESS OF THE				0				
						0				
10-	Fe staining below 10'									
						0				
	Red and gray clay, moist, stiff									
15						0				
						0				
20-										ı
								-		
	sand seams- wet									1
\exists										
25	End Of Boring ● 24' GW sample collected			£	<u> </u>	J.				
ᅴ										
\exists										
ᅴ										
30-										
во	RING DESIGNATION B-3	ASSOC	T C		C.	*** ***********************************		F		NUMBER OF 1

APPENDIX B

LABORATORY REPORTS AND CHAIN-OF-CUSTODY DOCUMENTATION



ANALYTICAL REPORT

JOB NUMBER: 318054 Project ID: 9003 N. MAIN

Prepared For:

ATC Associates, Inc. 3928 Bluebonnet Drive Stafford, TX 77477

Attention: Patrick Dworaczyk

Date: 07/10/2006

Signature

Name: Dean A. Joiner

Title: Project Manager II

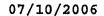
E-Mail: djoiner@stl-inc.com

Severn Trent Laboratories

6310 Rothway Drive Houston, TX 77040

PHONE: 713-690-4444

TOTAL NO. OF PAGES 48





Patrick Dworaczyk
ATC Associates, Inc.
3928 Bluebonnet Drive
Stafford, TX 77477

Reference:

Project : 9003 N. MAIN

Project No. : 318054
Date Received : 06/21/2006
STL Job : 318054

Dear Patrick Dworaczyk:

Enclosed are the analytical results for your project referenced above. The following samples are included in the report.

- 1. B-1 12-14
- 2. B-2 20-22
- 3. B-3 20-22
- 4. B-1
- 5. B-2
- 6. B-3
- 7. TRIP BLANK

All hold times were met for the tests performed on these samples.

Enclosed, please find the Quality Control Summary. All quality control results for the QC batch that are applicable to the sample(s) are acceptable except as noted in the QC batch reports.

The test results in this report meet all NELAP requirements for STL Houston's NELAP accredited parameters. Any exceptions to NELAP requirements will be noted and included in a case narrative as a part of this report.

If the report is acceptable, please approve the enclosed invoice and forward it for payment.

Thank you for selecting Severn-Trent Laboratories to serve as your analytical laboratory on this project. If you have any questions concerning these results, please feel free to contact me at any time.

We look forward to working with you on future projects.

Singerely,

Dean A. Joiner Project Manager



Table 1

Cross-Reference Field Sample Identifications and Laboratory Identifications

Field Identification	Laboratory Identification	8021B	TX1005	8315	Comment
B-1 12-14	318054-1	X	X	X	
B-2 20-22	318054-2	X	X	×	
B-3 20-22	318054-3	X	X	×	
B-1	318054-4	X	X	×	
B-2	318054-5	X	X	×	
B-3	318054-6	X	X	×	
TRIP BLANK	318054-7				Trip Blank; Not on C-O-C; No Tests Assigned

Appendix A Laboratory Data Package Cover Page

This data package consists of:

- This signature page, the laboratory review checklist, and the following reportable data:
- R1 Field chain-of-custody documentation;
- Sample identification cross-reference;
- R3 Test reports (analytical data sheets) for each environmental sample that includes:
 - a) Items consistent with NELAC 5.13 or ISO/IEC 17025 Section 5.10
 - b) dilution factors,
 - c) preparation methods,
 - d) cleanup methods, and
 - e) if required for the project, tentatively identified compounds (TICs).
- R4 Surrogate recovery data including:
 - a) Calculated recovery (%R), and
 - b) The laboratory's surrogate QC limits.
- Test reports/summary forms for blank samples; R5
- Test reports/summary forms for laboratory control samples (LCSs) including:
 - a) LCS spiking amounts,
 - b) Calculated %R for each analyte, and
 - c) The laboratory's LCS QC limits.
- Test reports for project matrix spike/matrix spike duplicates (MS/MSDs) including: R7
 - a) Samples associated with the MS/MSD clearly identified,
 - b) MS/MSD spiking amounts,
 - c) Concentration of each MS/MSD analyte measured in the parent and spiked samples,
 - d) Calculated %Rs and relative percent differences (RPDs), and
 - e) The laboratory's MS/MSD QC limits
- R8 Laboratory analytical duplicate (if applicable) recovery and precision:
 - a) the amount of analyte measured in the duplicate,
 - b) the calculated RPD, and
 - c) the laboratory's QC limits for analytical duplicates.
- R9 List of method quantitation limits (MQLs) for each analyte for each method and matrix;
- R10 Other problems or anomalies.
- The Exception Report for every "No" or "Not Reviewed (NR)" item in laboratory review checklist.

Release Statement: I am responsible for the release of this laboratory data package. This data package has been reviewed by the laboratory and is complete and technically compliant with the requirements of the methods used, except where noted by the laboratory in the attached exception reports. By me signature below, I affirm to the best of my knowledge, all problems/anomalies, observed by the laboratory as having the potential to affect the quality of the data, have been identified by the laboratory in the Laboratory Review Checklist, and no information or data have been knowingly withheld that would affect the quality of the data.

Check, if applica	able: [] This laborat	ory is an in-hous	se laboratory controlled by	y the p	person	la tha
4.	trong friends a der a la cionata	ng the cover pag	e of the rule-required repo	or an	n exan	pie, me z sionature
APAR) in w	hich these data are used	is responsible 10	r releasing this data packs	rgc an	,	, 516,11111
affirming th	e above release statement	t ip true.				
	Washington	Nhor	Laboratory Director	7	112	100
Norman Flynn	Composit	0	Official Title (printed)		Date	L
Name (Printed)	Signature	()				

Lab	orato	ry Name: STL-Houston LRC Da	te: 06/26/06					
			ory Job Number: 318054					
<u> </u>			tch Number(s): 157296 (Soil)-TX10	05			~~~~	
			tell Number(s). 137290 (30h)-17(10h	Yes	No	NIA3	NR ⁴	ED+
# ¹	A ²	Description		105	INO	IVA	INK	5
		Chain-of-custody (C-O-C)					4,10	
RI	OI	Did samples meet the laboratory's standard conditions of sample a	cceptability upon receipt?		X			1
		Were all departures from standard conditions described in an excep		X				
R2	OI	Sample and quality control (QC) identification				7.2		
		Are all field sample ID numbers cross-referenced to the laboratory	ID numbers?	X				
	1	Are all laboratory ID numbers cross-referenced to the corresponding		X				
R3	OI	Test reports						
		Were all samples prepared and analyzed within holding times?		X				
		Other than those results < MQL, were all other raw values bracket	ed by calibration standards?	X				
		Were calculations checked by a peer or supervisor?		X				
ļ		Were all analyte identifications checked by a peer or supervisor?		X				
		Were sample quantitation limits reported for all analytes not detect	ed?	X				
		Were all results for soil and sediment samples reported on a dry we		X				
	ĺ	Were % moisture (or solids) reported for all soil and sediment sam		X				
		If required for the project, TICs reported?				X		
R4	o	Surrogate recovery data						
	 _ _ _ _ _ _ _ _ _ _ _ _ _ 	Were surrogates added prior to extraction?		X				
	1	Were surrogate percent recoveries in all samples within the laborate	ory OC limits?	X				\vdash
R5	OI	Test reports/summary forms for blank samples	ory QC mmo.	 				
-	101	Were appropriate type(s) of blanks analyzed?		X				\vdash
l		Were blanks analyzed at the appropriate frequency?		X				-
		Were method blanks taken through the entire analytical process, in	cluding prepa/ation and if	X				
		applicable, cleanup procedures?	cidding preparation and, it	1				
		Were blank concentrations < MQL?		X				一
R6	OI	Laboratory control samples (LCS):			 -			
100	<u> </u>	Were all COCs included in the LCS?		-	X			2
l	1	Was each LCS taken through the entire analytical procedure, inclu	ling area and cleanum stems?	X	1			F
		Were LCSs analyzed at the required frequency?	ing prep and eleurop steps:	X				
	1	Were LCS (and LCSD, if applicable) %Rs within the laboratory Q	? limite?	X				
	1	Does the detectability data document the laboratory's capability to		Λ		X		-
		to calculate the SQLs?	detect the COCs at the MDL used			^		
İ		Was the LCSD RPD within QC limits?		X	-			
R7	01	Matrix spike (MS) and matrix spike duplicate (MSD) data		^				
<u> </u>	101	Were the project/method specified analytes included in the MS and	MSD2	х				-
		Were MS/MSD analyzed at the appropriate frequency?	INOD:	X				
İ		Were MS (and MSD, if applicable) %Rs within the laboratory QC	limite?	<u>^</u>	-		X	3
1	l	Were MS/MSD RPDs within laboratory QC limits?	mms:	-			X	3
R8	OI	Analytical duplicate data			-			<u> </u>
No.	101	Were appropriate analytical duplicates analyzed for each matrix?				x		
		Were analytical duplicates analyzed at the appropriate frequency?				$\frac{\lambda}{X}$		-
			? limito?			$\hat{\mathbf{x}}$		-
-	-	Were RPDs or relative standard deviations within the laboratory Q	C mms;					
R9	OI	Method quantitation limits (MQLs):	inte makaga?	X				
1		Are the MQLs for each method analyte included in the laboratory		X				
}		Do the MQLs correspond to the concentration of the lowest non-zer	to campianon standaro?	X				-
D-10	<u></u>	Are unadjusted MQLs included in the laboratory data package?		^				
R10	OI	Other problems/anomalies	T DC and ED9	х			-	
l		Are all known problems/anomalies/special conditions noted in this		X				-
		Were all necessary corrective actions performed for the reported day. Was applicable and available technology used to lower the SQL to		X	-			-
		affects on the sample results?	minimize me manta interference	^				L
		<u></u>						

SX

- Items identified by the letter "R" must be included in the laboratory data package submitted in the TRRP-required report(s). Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.
 O= organic analyses; I = inorganic analyses (and general chemistry, when applicable);
- NA = Not applicable;
- NR = Not reviewed;
- ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

ΑI

Ap	pen	dix A (cont'd): Laboratory Review Checklis	st: Reportable Data					
Lab	orato	ry Name: STL-Houston	LRC Date: 06/26/06					
Proj	ect N	Iame: 9003 N. MAIN	Laboratory Job Number: 318054					
Rev	iewe	r Name: MW	Prep Batch Number(s): 157296 (Soil)-TX	1005				
#1	A ²	Description		Yes	No	NA3	NR ⁴	ER#5
S1		Initial calibration (ICAL)		1111			25.00	
~		Were response factors and/or relative response factors for each	h analyte within OC limits?	X		†	†	
		Were percent RSDs or correlation coefficient criteria met?	<u> </u>	X		 		
		Was the number of standards recommended in the method use	ed for all analytes?	X				
		Were all points generated between the lowest and highest stan		X				
		Are ICAL data available for all instruments used?		X			T	
		Has the initial calibration curve been verified using an approp	riate second source standard?	X	 	 	 	1
S2	OI	Initial and continuing calibration verification (ICCV and		1	1.10	1		
-	<u>~</u> _	Was the CCV analyzed at the method-required frequency?		X			1	1
		Were percent differences for each analyte within the method-	required OC limits?	X	 			†
		Was the ICAL curve verified for each analyte?		$\frac{\hat{\mathbf{x}}}{\mathbf{x}}$		 	 	
		Was the absolute value of the analyte concentration in the inor	rganic CCB < MDL?	† ^	 	X	 	
S3	0	Mass spectral tuning:	PARALLE STATE STATE AND ADDRESS OF THE PARALLES AND ADDRES	1	100	 ``	1 . 1	1.74
22		Was the appropriate compound for the method used for tuning	79	 	 	X	1	1
		Were ion abundance data within the method-required QC limit		 	 	X	 	
S4	0	Internal standards (IS):	ω:	 	 	1	 	+
34	<u> </u>	Were IS area counts and retention times within the method-red	united OC limite?	├		X		
OE	OI.	Raw data (NELAC section 1 appendix A glossary, and section 1	tion 5 12 on ISO/ISC 17025 section	-	 	<u> </u>	 	
S5	<u>OI</u>	Were the raw data (for example, chromatograms, spectral data		Х		├	 	
				$\frac{\hat{x}}{x}$	├	 	 	
06		Were data associated with manual integrations flagged on the	raw datar	 ^-				
S6	<u>o</u>	Dual column confirmation	-1009	-		X		
		Did dual column confirmation results meet the method-require	ea QC?			A	17.57	-
S7	<u>o_</u>	Tentatively identified compounds (TICs):	1:					
		If TICs were requested, were the mass spectra and TIC data su	ibject to appropriate checks?		11 2 3	X	1 1 2 3 3 3 1 1	
S8	1	Interference Check Sample (ICS) results:				-		1
		Were percent recoveries within method QC limits?		-	7.7.	X		-
S9	<u> </u>	Serial dilutions, post digestion spikes, and method of stand	ard additions				1 2 2 2	
		Were percent differences, recoveries, and the linearity within	the QC limits specified in the method?	!	ļ	X		
S10	<u>01</u>	Method detection limit (MDL) studies		ļ				
		Was a MDL study performed for each reported analyte?		X	 	ļ	<u> </u>	
		Is the MDL either adjusted or supported by the analysis of DC	Ss?	X		ļ	ļ	
S11	<u>OI</u>	Proficiency test reports:		<u> </u>	<u> </u>	 	<u> </u>	ļ
		Was the laboratory's performance acceptable on the applicable	e proficiency tests or evaluation studies?	X	<u> </u>			
S12	OI	Standards documentation			ļ	<u> </u>		1
		Are all standards used in the analyses NIST-traceable or obtain	ned from other appropriate sources?	X	<u> </u>			<u> </u>
S13	OI	Compound/analyte identification procedures						
		Are the procedures for compound/analyte identification documents	nented?	X	<u> </u>			<u> </u>
S14	OI	Demonstration of analyst competency (DOC)					1. 1.50	
		Was DOC conducted consistent with NELAC Chapter 5C or I		X				<u> </u>
j	-	Is documentation of the analyst's competency up-to-date and	on file?	X				
S15	OI	Verification/validation documentation for methods (NELA	C Chap 5 or ISO/IEC 17025 Section 5)	17.00	35, 3	7,45%		13174
		Are all the methods used to generate the data documented, ver	ified, and validated, where applicable?	X				
	~~	Laboratory standard operating procedures (SOPs):					1700	10.000
S16	OI							

Items identified by the letter "R" should be included in the laboratory data package submitted to the TCEQ in the TRRP-required report(s). Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.

O = organic analyses; I = inorganic analyses (and general chemistry, when applicable).

5

NA = Not applicable.

NR = Not Reviewed.

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

Appe	ndix A (cont'd): Laboratory Review Cho	ecklist: Exception Reports				
Labora	itory Name: STL-Houston	LRC Date: 06/26/06				
Project	Name: 9003 N. MAIN	Laboratory Job Number: 318054				
Reviev	ver Name: MW	Prep Batch Number(s): 157296 (Soil)-TX1005				
ER#	DESCRIPTION					
1	The temperature of the cooler received by the laboratory on 06/21/06 was above the acceptable range of 2.0-6.0 °C.					
2	The final concentration of any hydrocarbons d	required by the method, this range was not spiked into the LCS/LCSD. letected in this range was calculated from the response factor of the C12-raction efficiency of the C28-C35 range hydrocarbons was determined ns.				
3	The laboratory selected another client's sample	e to perform as the MS/MSD.				

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked on the LRC)

·		ry Name: STL-Houston LRC D	ate: 06/26/06				-	
			tory Job Number: 318054					
		· · · · · · · · · · · · · · · · · · ·				····		
Rev	iewer	r Name: MW Prep B	atch Number(s): 157449 (Water)-TXI					
# ¹	A ²	Description		Yes	No	NA ³	NR ⁴	ER#
		Chain-of-custody (C-O-C)			A			110011
R1	OI		cceptability upon receipt?	 	X			1
		Were all departures from standard conditions described in an exce		X				
R2	OI	Sample and quality control (QC) identification		-				
	-	Are all field sample ID numbers cross-referenced to the laboratory	ID numbers?	X				
	1	Are all laboratory ID numbers cross-referenced to the correspondi		X				
R3	OI	Test reports		 		-		
	 	Were all samples prepared and analyzed within holding times?		X				
		Other than those results < MQL, were all other raw values bracket	ed by calibration standards?	$\frac{\Lambda}{X}$				
	1	Were calculations checked by a peer or supervisor?	od by cantraction standards:	X	 			
		Were all analyte identifications checked by a peer or supervisor?		$\frac{\Lambda}{X}$				
		Were sample quantitation limits reported for all analytes not detec	ted?	X				
		Were all results for soil and sediment samples reported on a dry w			-	x		├
		Were % moisture (or solids) reported for all soil and sediment sam				X		
l	1		pies:			X		
R4	0	If required for the project, TICs reported?						
K4	<u> </u>	Surrogate recovery data						
		Were surrogates added prior to extraction?	001, 70	X				
200		Were surrogate percent recoveries in all samples within the labora	tory QC limits?	X				ļ
R5	OI	Test reports/summary forms for blank samples			ļ			
		Were appropriate type(s) of blanks analyzed?		X				ļ
		Were blanks analyzed at the appropriate frequency?		X	ļ			
		Were method blanks taken through the entire analytical process, in	icluding prepa/ation and, if	X				İ
l		applicable, cleanup procedures?		-,,				ļ
		Were blank concentrations < MQL?		X		-		 -
R6	OI	Laboratory control samples (LCS):						
		Were all COCs included in the LCS?			Х			2
	1	Was each LCS taken through the entire analytical procedure, inclu	ding prep and cleanup steps?	X				ļ
		Were LCSs analyzed at the required frequency?		X				ļ
		Were LCS (and LCSD, if applicable) %Rs within the laboratory Q		X				ļ
		Does the detectability data document the laboratory's capability to	detect the COCs at the MDL used			X		ĺ
		to calculate the SQLs?						ļ
		Was the LCSD RPD within QC limits?		X				<u> </u>
R7	OI	Matrix spike (MS) and matrix spike duplicate (MSD) data						ļ
		Were the project/method specified analytes included in the MS an	d MSD?	X				ļ
		Were MS/MSD analyzed at the appropriate frequency?		X				ļ
		Were MS (and MSD, if applicable) %Rs within the laboratory QC	limits?	X				ļ
	<u> </u>	Were MS/MSD RPDs within laboratory QC limits?		X				
R8	OI	Analytical duplicate data						ļ
		Were appropriate analytical duplicates analyzed for each matrix?				X		
		Were analytical duplicates analyzed at the appropriate frequency?				X		
		Were RPDs or relative standard deviations within the laboratory Q	C limits?			Х		Ĺ
R9	OI	Method quantitation limits (MQLs):						
		Are the MQLs for each method analyte included in the laboratory		X				
		Do the MQLs correspond to the concentration of the lowest non-ze	ero calibration standard?	X				
		Are unadjusted MQLs included in the laboratory data package?		Х				
R10	OI	Other problems/anomalies						
		Are all known problems/anomalies/special conditions noted in this	LRC and ER?	Х				
	1	Were all necessary corrective actions performed for the reported d	ata?	X				
		Was applicable and available technology used to lower the SQL to		X				
	l	affects on the sample results?						

SX

- Items identified by the letter "R" must be included in the laboratory data package submitted in the TRRP-required report(s). Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.
 O= organic analyses; I = inorganic analyses (and general chemistry, when applicable);
- NA = Not applicable; NR = Not reviewed;
- ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

Αl

Reviewer Name	Name: STL-Houston	LRC Date: 06/26/06					
#	e: 9003 N. MAIN	Laboratory Job Number: 318054					
# A Desc S1 OI Initia	me: MW	Prep Batch Number(s): 157449 (Water)-7	'X10)5	····	*******	
S1 OI Initial Were Was Were Was Was Was Was Was Was Were Was Were Were Were Were Were Were S6 O Dual Did d S7 O Tental Tental Tental Tental Tental Tental Tental Were S9 I Seria Were S10 OI Meth Was a Is the S11 OI Profi Was a Is the S12 OI Stand Are a S13 OI Comp Are till S14 OI Demot Was a Is doc S15 OI Verif S15 OI OI OI OI OI OI OI O		Top Sates Trained (a), 15 This (Trates)					ER#
Were Were Was Were	tial calibration (ICAL)		103	110	INA	HAX.	Lin
Were Was Were	ere response factors and/or relative response factors for each	h analyta within OC limita?	X		 		
Was Were	ere percent RSDs or correlation coefficient criteria met?	n analyte within QC hmits?	X		 		
Were Are I	as the number of standards recommended in the method use	ad for all analytes?	X			 	
Are I	ere all points generated between the lowest and highest star		X		 	├	
Has t S2 OI Initia Was Was Was Was Was Was Were Were Were Were Were S5 OI Raw Were Were S6 O Dual Did d S7 O Tenta If TIC S8 I Inter Were S9 I Seria Were S11 OI Meth Was a Is the S12 OI Stanc Are a S13 OI Comp Are till S14 OI Demot Was I S6 OI Verif Was I S15 OI Verif Was I S15 OI Verif	e ICAL data available for all instruments used?	idard used to carculate the curve?	X		-	 	
S2 OI Initia Was Were Was Were	s the initial calibration curve been verified using an approp	riote second source standard?	X		 		
Was Were Was Were Was Were	tial and continuing calibration verification (ICCV and (^	66.1			
Were Was	is the CCV analyzed at the method-required frequency?	CCV) and continuing candiation	X			-	-
Was Was	ere percent differences for each analyte within the method-	required OC limite?	X				
Was Was	is the ICAL curve verified for each analyte?	required QC minits:	X		 		
S3 O Mass Were	is the real curve verified for each analyte? Is the absolute value of the analyte concentration in the ino	roanic CCR < MDI 2	_^_		X	-	
Was Were	is the absolute value of the analyte concentration in the mo	Iganic CCB < WDL?			^	10.00	35.5
Were	is the appropriate compound for the method used for tuning				X		
S4 O Inter	ere ion abundance data within the method-required QC limi				X		
Were Were	ernal standards (IS):	118 ;					
S5 OI Raw Were	ernal standards (18): ere IS area counts and retention times within the method-rec	avirad OC limite?	ļ		X		
Were Were	w data (NELAC section 1 appendix A glossary, and sec						
Were	ere the raw data (for example, chromatograms, spectral data		X			-	
S6 O Dual	ere data associated with manual integrations flagged on the		X				<u> </u>
Did d S7 O Tents If TiC S8 I Inter Were S9 I Seria Were S10 OI Meth Was a Is the S11 OI Profit Was 1 S12 OI Stance Are a S13 OI Com Are tit S14 OI Demo Was 1 Is doc S15 OI Verif	al column confirmation	Taw data?	-		. 973.		
S7 O Tentic	I dual column confirmation results meet the method-require	MOC2			X		
If TIC S8 I Inter Were S9 I Seria Were S10 OI Meth Was a Is the S11 OI Profit Was 1 S12 OI Stance Are a S13 OI Com Are the S14 OI Demo Was 1 Is doc S15 OI Verif		ed QC;					
S8 I Inter	ntatively identified compounds (TICs): TICs were requested, were the mass spectra and TIC data so	which to appropriate shocked			X	31/25/10	
Were Were	erference Check Sample (ICS) results:	doject to appropriate checks?	9 (%)	in iga.	-^-	0.135.5	111
Seria Seria	errefrence Check Sample (1CS) results: ere percent recoveries within method QC limits?				Х		
Were Were			GgSA.				10,00
S10 OI Meth Was a Is the	rial dilutions, post digestion spikes, and method of stand are percent differences, recoveries, and the linearity within				Х		
Was a Is the	thod detection limit (MDL) studies	the QC mints specified in the method?	72.5	71.75	_^_		
Is the	is a MDL study performed for each reported analyte?		х				
S11 OI	he MDL either adjusted or supported by the analysis of DC	200	$\frac{\lambda}{X}$				
Was t S12 OI Stand Are a S13 OI Com Are tt S14 OI Demo Was I Is doc	oficiency test reports:	, JS :	^		1 77 1		
S12 OI Stand Are a	is the laboratory's performance acceptable on the applicable	e proficiency tests or evaluation studies?	Х				
Are a	ndards documentation	e protected y tests of evaluation studies?	<u>^</u>	77.7	-	1 12 1	
S13 OI Com Are th S14 OI Demo Was I Is doc S15 OI Verif	all standards used in the analyses NIST-traceable or obtain	ned from other engraprists courses?	Х				
Are the S14 OI Demo Was I Is doc S15 OI Verif	mpound/analyte identification procedures	ned from other appropriate sources:	^	75.7	5.7 (4	1,75,175	Jan 12, 1
Name	the procedures for compound/analyte identification documents	nantad?	X				
Was I Is doc S15 OI Verif		AVIIIO I	7			3355	: दिवसी र
Is doc S15 OI Verif	monstration of analyst competency (DOC) s DOC conducted consistent with NELAC Chapter 5C or I	SO/IEC 42	\mathbf{x}				
S15 OI Verif	s DOC conducted consistent with NEEAC Chapter 3C of I		$\frac{\hat{\mathbf{x}}}{\mathbf{x}}$				
AT A CLI	rification/validation documentation for methods (NELA			199	127	4,55	VA.
Aran	e all the methods used to generate the data documented, ver	rified and validated where applicable?	Х				V17411
		ined, and vanuated, where applicable?					7 - 4
S16 OI Labo	boratory standard operating procedures (SOPs): aboratory SOPs current and on file for each method performs.		x				

Items identified by the letter "R" should be included in the laboratory data package submitted to the TCEQ in the TRRP-required report(s).

Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.

O = organic analyses; I = inorganic analyses (and general chemistry, when applicable).

NA = Not applicable.

NR = Not Reviewed.

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

Appendix A (cont'd): Laboratory Review Checklist: Exception Reports							
Labora	tory Name: STL-Houston	LRC Date: 06/26/06					
Project Name: 9003 N. MAIN		Laboratory Job Number: 318054					
Reviewer Name: MW		Prep Batch Number(s): 157449 (Water)-TX1005					
ER#1	DESCRIPTION						
1	The temperature of the cooler received by the	laboratory on 06/21/06 was above the acceptable range of 2.0-6.0 °C.					
2		required by the method, this range was not spiked into the LCS/LCSD. etected in this range was calculated from the response factor of the C12-					
		raction efficiency of the C28-C35 range hydrocarbons was determined					

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked on the LRC)

Lab	orato	ry Name: STL-Houston LR	C Date: 06/26/06					
			poratory Job Number: 318054					
<u> </u>				r				
#I			p Batch Number(s): 157581 (Soil)-BTEX		λ1-	NIA 3	NID 4	1215.41
#	A	Description Co. Co. Co. Co. Co. Co. Co. Co. Co. Co.		Yes	No	INA	INK	ER#
nı	01	Chain-of-custody (C-O-C)				59,5		(core)
R1	U	Did samples meet the laboratory's standard conditions of samp		 	X	ļ		1
	-	Were all departures from standard conditions described in an e	exception report?	X	ļ			
R2	OI	Sample and quality control (QC) identification						111
		Are all field sample ID numbers cross-referenced to the labora		X		ļ		
72.0	-	Are all laboratory ID numbers cross-referenced to the correspondence	onding QC data?	X			X	
R3	OI	Test reports				ļ		
		Were all samples prepared and analyzed within holding times		X	ļ	ļ		ļ
		Other than those results < MQL, were all other raw values bra-	cketed by calibration standards?	X		ļ		ļ
		Were calculations checked by a peer or supervisor?		X		X		
		Were all analyte identifications checked by a peer or supervisor		X	ļ			
		Were sample quantitation limits reported for all analytes not d		X			 	
		Were all results for soil and sediment samples reported on a dr		X				
	1	Were % moisture (or solids) reported for all soil and sediment	samples?	X				
	<u> </u>	If required for the project, TICs reported?				X		
R4	0	Surrogate recovery data						
		Were surrogates added prior to extraction?		X				
	<u> </u>	Were surrogate percent recoveries in all samples within the lab	oratory QC limits?		X			2
R5	OI	Test reports/summary forms for blank samples						
		Were appropriate type(s) of blanks analyzed?		X				
		Were blanks analyzed at the appropriate frequency?		X				
		Were method blanks taken through the entire analytical proces	s, including preparation and, if	X				
		applicable, cleanup procedures?						
		Were blank concentrations < MQL?		X				
R6	OI	Laboratory control samples (LCS):						
		Were all COCs included in the LCS?		X				
		Was each LCS taken through the entire analytical procedure, in	ncluding prep and cleanup steps?	X				
		Were LCSs analyzed at the required frequency?		X				~~~~
		Were LCS (and LCSD, if applicable) %Rs within the laborator	y QC limits?	X				
		Does the detectability data document the laboratory's capabilit	y to detect the COCs at the MDL used			X		
		to calculate the SQLs?						
		Was the LCSD RPD within QC limits?				X		
R7	OI	Matrix spike (MS) and matrix spike duplicate (MSD) data						
		Were the project/method specified analytes included in the MS	and MSD?	Х				***************************************
		Were MS/MSD analyzed at the appropriate frequency?		X				
		Were MS (and MSD, if applicable) %Rs within the laboratory	OC limits?	х				
		Were MS/MSD RPDs within laboratory QC limits?	·	Х	$\neg \neg$			
R8	OI	Analytical duplicate data						
		Were appropriate analytical duplicates analyzed for each matri	x?			X		
		Were analytical duplicates analyzed at the appropriate frequen				X		
		Were RPDs or relative standard deviations within the laborator				X		
R9	OI	Method quantitation limits (MQLs):						
	<u> </u>	Are the MQLs for each method analyte included in the laborate	orv data package?	X			\neg	
		Do the MQLs correspond to the concentration of the lowest no		$\frac{x}{x}$				
		Are unadjusted MQLs included in the laboratory data package		$\frac{\Lambda}{X}$				
R10	OI	Other problems/anomalies						
	"	Are all known problems/anomalies/special conditions noted in	this I RC and FR?	X				
		Were all necessary corrective actions performed for the reporte		$\frac{\hat{X}}{X}$				
		Was applicable and available technology used to lower the SQ		$\frac{\Delta}{X}$				
		affects on the sample results?	a to minimize the matrix interference	^	ļ		1	
	Ļ	Items identified by the letter "P" must be included in the laboratory data	The state of the s	لببا			!	

^{1.} Items identified by the letter "R" must be included in the laboratory data package submitted in the TRRP-required report(s). Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.

2. = organic analyses; I = inorganic analyses (and general chemistry, when applicable);

3. NA = Not applicable;

4. NR = Not reviewed;

5. ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

Lab	orato	ry Name: STL-Houston LR	C Date: 06/26/06					
Proj	ect N	lame: 9003 N. MAIN La	boratory Job Number: 318054					
Rev	iewei	r Name: ERA	rep Batch Number(s): 157581 (Soil)-BTEX			,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		
#1		Description		Yes	No	NA ³	NR ⁴	ER#
					77.5	9 11 1	77.5	
<u>S1</u>	OI	Initial calibration (ICAL) Were response factors and/or relative response factors for each	analyta within OC limits?	X			 	
		Were percent RSDs or correlation coefficient criteria met?	anaryte within OC mints:	X		 		
		Was the number of standards recommended in the method used	for all analytee?	X		 	 	
		Were all points generated between the lowest and highest stand	lord used to calculate the curve?	X			 	
			lard used to calculate the curve:	X	-	 	 	\vdash
		Are ICAL data available for all instruments used? Has the initial calibration curve been verified using an appropri	isto second source standard?	X	 	 	 	
~	2.0	Initial and continuing calibration verification (ICCV and C	CV) and continuing calibration	<u> </u>				
S2	OI		CV) and continuing campi atton	X		 	 	\vdash
		Was the CCV analyzed at the method-required frequency? Were percent differences for each analyte within the method-re	equired OC limite?	X	\vdash	 	 	t
		were percent differences for each analyte within the method-re	Adired AC illinist	$\frac{\Lambda}{X}$	 	NA3 NR4 F	 	
		Was the ICAL curve verified for each analyte? Was the absolute value of the analyte concentration in the inorg	ronio CCR < MDI 2	<u> </u>	 	Y	 	1-
~~			gaine CCB < MDL!	20127	-	1-	-	
S3	0	Mass spectral tuning:	3		-	V	 	
		Was the appropriate compound for the method used for tuning		-			 	
~ -		Were ion abundance data within the method-required QC limits	S?		-	<u> </u>	 	1
<u>54</u>	0	Internal standards (IS):	-in-1 OC limite?	X				┼──
		Were IS area counts and retention times within the method-req	f 12 YCO/TEC 17025		-	-	 	┼
S5_	OI	Raw data (NELAC section 1 appendix A glossary, and section	on 5.12 or 180/IEC 1/025 section	X			-	┼─
		Were the raw data (for example, chromatograms, spectral data)		X		 	┼	┼
		Were data associated with manual integrations flagged on the r	aw data?	<u> </u>	-	-	 	┼──
S6	0	Dual column confirmation	1000	X			-	┼
		Did dual column confirmation results meet the method-required	a QC?	1		 	1 1111	┼─
<u>S7</u>	0	Tentatively identified compounds (TICs):				37	-	-
		If TICs were requested, were the mass spectra and TIC data su	bject to appropriate checks?		1, 1		-	+
S8	I	Interference Check Sample (ICS) results:		71, 27				-
		Were percent recoveries within method QC limits?			1000		1	+
<u>59</u>	I	Serial dilutions, post digestion spikes, and method of standa	ard additions			1	-	
	L	Were percent differences, recoveries, and the linearity within the	he QC limits specified in the method?		11.	<u>^</u>	 	┼
S10	OI	Method detection limit (MDL) studies		-		 		┼—
	<u> </u>	Was a MDL study performed for each reported analyte?		X		 	 	┼
		Is the MDL either adjusted or supported by the analysis of DC	Ss?	X		 	-	
S11	OI	Proficiency test reports:		125			 	┼
		Was the laboratory's performance acceptable on the applicable	proficiency tests or evaluation studies?	X		ļ	 	┼
S12	OI	Standards documentation		-		ļ	 	-
		Are all standards used in the analyses NIST-traceable or obtain	ed from other appropriate sources?	X		ļ	 	
S13	OI	Compound/analyte identification procedures				<u> </u>	-	-
		Are the procedures for compound/analyte identification docum	ented?	X	ļ	-	 	╀
S14	OI	Demonstration of analyst competency (DOC)				ļ	1-	4
		Was DOC conducted consistent with NELAC Chapter 5C or IS	SO/IEC 4?	X	 	<u> </u>	├ ─	₩
	<u></u>	Is documentation of the analyst's competency up-to-date and o	n file?	X		ļ		—
S15	OI	Verification/validation documentation for methods (NELAC	C Chap 5 or ISO/IEC 17025 Section 5)	35.1				1
	1	Are all the methods used to generate the data documented, veri	fied, and validated, where applicable?	X				
S16	OI	Laboratory standard operating procedures (SOPs):					1 1 1 1 1	
	 ~~	Are laboratory SOPs current and on file for each method perform	rmed?	X			1	

Items identified by the letter "R" should be included in the laboratory data package submitted to the TCEQ in the TRRP-required report(s). Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.

O = organic analyses; I = inorganic analyses (and general chemistry, when applicable).

NA = Not applicable.

NR = Not Reviewed.

³

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

Apper	ndix A (cont'd): Laboratory Revie	w Checklist: Exception Reports			
Laborat	ory Name: STL-Houston	LRC Date: 06/26/06			
Project	Name: 9003 N. MAIN	Laboratory Job Number: 318054			
Review	er Name: ERA	Prep Batch Number(s): 157581 (Soil)-BTEX			
ER#1	DESCRIPTION				
1	The temperature of the cooler received	by the laboratory on 06/21/06 was above the acceptable range of 2.0-6.0 °C.			
2		The a,a,a-trifluorotoluene surrogate recoveries on both columns and the bromofluorobenzene recovery on column SPB-624 in sample 318054-2 were outside acceptance limits due to matrix interference.			

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked on the LRC)

Lab	orato	ry Name: STL-Houston LR	RC Date: 06/26/06					
		· · · · · · · · · · · · · · · · · · ·	boratory Job Number: 318054					
				T237				
			ep Batch Number(s): 15/626 (water)-B1.			120.3	T A	
#.	A.			Yes	No	NA'	NR"	ER#5
	~-							315 7
RI	OI	Did samples meet the laboratory's standard conditions of sam	ple acceptability upon receipt?	ļ	X	<u> </u>		1
	ļ		exception report?	X				
Laboratory Name: STL-Houston								
	Aboratory Name: STL-Houston Project Name: 9003 N. MAIN Reviewer Name: BRA #I A Description Chain-of-custody (C-O-C) Did samples meet the laboratory's standard conditions of Were all departures from standard conditions described Are all field sample ID numbers cross-referenced to the Are all laboratory ID numbers cross-referenced to the Are all laboratory ID numbers cross-referenced to the Are all laboratory ID numbers cross-referenced to the Are all laboratory ID numbers cross-referenced to the Are all laboratory ID numbers cross-referenced to the Are all laboratory ID numbers cross-referenced to the Are all laboratory ID numbers cross-referenced to the Are all laboratory ID numbers cross-referenced to the Are all laboratory ID numbers cross-referenced to the Are all laboratory ID numbers cross-referenced to the Are all laboratory iD numbers cross-referenced to the Are all laboratory iD numbers cross-referenced to the Are all laboratory exited the Are all laboratory and the Are all laboratory control analyzed Were all analyte identifications checked by a peer or supervisor? Were all nalyte identifications checked by a peer or supervisor? Were all results for soil and sediment samples reported to Were % moisture (or solids) reported for all soil and sed If required for the project, TICs reported? Were surrogates added prior to extraction? Were surrogates added prior to extraction? Were surrogate percent recoveries in all samples within Were blanks analyzed at the appropriate frequency? Were blanks analyzed at the appropriate frequency? Were blanks analyzed at the required frequency? Were all COCs included in the LCS? Was each LCS taken through the entire analytical procedures analyted the SQLs? Was the LCSD RPD within QC limits? OI Matrix spike (MS) and matrix spike duplicate (MSD) Were MS/MSD analyzed at the appropriate frequency? Were MS/MSD analyzed at the appropriate frequency? Were MS (and MSD, if applicable) %Rs within the laboratory were MS/MSD RPDs within laboratory QC limits? Are the M							
			onding QC data?	X		<u></u>	<u> </u>	
R3	101			 				
					<u> </u>			
			acketed by calibration standards?		 	ļ		ļ
	İ		analyzed within holding times? A were all other raw values bracketed by calibration standards? A peer or supervisor? A peer or supervi					
					<u> </u>	ļ		
						ļ		
						<u> </u>		
			t samples?	X		L		
	<u> </u>					X		
R4	0						X	
				X				
			boratory QC limits?		X			2
R5 (OI	Test reports/summary forms for blank samples						
		Were blanks analyzed at the appropriate frequency?						
			ss, including preparation and, if	X				
				X				
R6	01							
			including prep and cleanup steps?					
							X	
				X				
			ty to detect the COCs at the MDL used			X	X	
				\sqcup				
	~=					X		
R7	OI_							
			S and MSD?			X		
				X				
			QC limits?					3
				 			X	3
К8	OI_							
			· · · · · · · · · · · · · · · · · · ·	lacksquare		-		
			ry QC limits?			X		
K9	<u>01</u>		MMMV - Magnet - 1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-	ليا				
ļ							$-\!\!\perp$	
			37	X				
KIU	OI	}		لييا				
								
			L to minimize the matrix interference	X	- 1		1	
		affects on the sample results?						

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A	per	ndix A (cont'd): Laboratory Review Checkli	st: Reportable Data					
Lat	orato	ory Name: STL-Houston	RC Date: 06/26/06					
Pro	ject N	Name: 9003 N. MAIN	aboratory Job Number: 318054					
Rev	iewe	r Name: ERA	Prep Batch Number(s): 157626 (Water)-B	TEX			,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	
#1	A ²	Description		Yes	No	NA ³	NR ⁴	ER#5
S1	01	Initial calibration (ICAL)						77.0
	1	Were response factors and/or relative response factors for each	th analyte within OC limits?	X	1			
	1	Were percent RSDs or correlation coefficient criteria met?		X				1
		Was the number of standards recommended in the method us	ed for all analytes?	X				
		Were all points generated between the lowest and highest star		X				1
		Are ICAL data available for all instruments used?		X				
		Has the initial calibration curve been verified using an approp	oriate second source standard?	X				
S2	OI	Initial and continuing calibration verification (ICCV and						
		Was the CCV analyzed at the method-required frequency?		X				
		Were percent differences for each analyte within the method-	required QC limits?	X	1			
		Was the ICAL curve verified for each analyte?		X	T			
		Was the absolute value of the analyte concentration in the inc	organic CCB < MDL?		Ī	X		
S3	0	Mass spectral tuning:						
		Was the appropriate compound for the method used for tuning	g?			X		
		Were ion abundance data within the method-required QC lim			1	X		
S4	0	Internal standards (IS):		1.4			10 10 1	
		Were IS area counts and retention times within the method-re	quired QC limits?	X				
S5	OI	Raw data (NELAC section 1 appendix A glossary, and sec	tion 5.12 or ISO/IEC 17025 section		1 1 14		7 T Z	
		Were the raw data (for example, chromatograms, spectral data	a) reviewed by an analyst?	X				
		Were data associated with manual integrations flagged on the	raw data?	X				
S6	0	Dual column confirmation			1	4.474	1111	100
		Did dual column confirmation results meet the method-requir	ed QC?		X			4
S7	0	Tentatively identified compounds (TICs):						
		If TICs were requested, were the mass spectra and TIC data s	ubject to appropriate checks?			X		
S8	I	Interference Check Sample (ICS) results:		1 7 7	1 5,21 5,	177	N 10.	
		Were percent recoveries within method QC limits?				X		
S9	I	Serial dilutions, post digestion spikes, and method of stand	lard additions	1 1 1 1 1 1	19.35	44	1.0 4 1.1	
		Were percent differences, recoveries, and the linearity within	the QC limits specified in the method?			X		
S10	OI	Method detection limit (MDL) studies		151.5	10 Sec.	1.5	134.12	
		Was a MDL study performed for each reported analyte?		X				
		Is the MDL either adjusted or supported by the analysis of DC	CSs?	X				
S11	OI	Proficiency test reports:		45.5	170.3	7.00	1.7	
		Was the laboratory's performance acceptable on the applicable	e proficiency tests or evaluation studies?	X			l	
S12	Ol	Standards documentation				1.41	ring .	1.3
		Are all standards used in the analyses NIST-traceable or obtain	ned from other appropriate sources?	X				
S13	OI	Compound/analyte identification procedures						
		Are the procedures for compound/analyte identification documents	mented?	X				
S14	OI	Demonstration of analyst competency (DOC)		3.7				<u> </u>
		Was DOC conducted consistent with NELAC Chapter 5C or l		X				
	<u> </u>	Is documentation of the analyst's competency up-to-date and		X				
S15	OI	Verification/validation documentation for methods (NELA						1.1 1
		Are all the methods used to generate the data documented, ver	rified, and validated, where applicable?	Х				
S16	OI	Laboratory standard operating procedures (SOPs):			1,14	77 1 14		117.5
		Are laboratory SOPs current and on file for each method perfe	ormed?	X				
	L			<u> </u>			L	L

Items identified by the letter "R" should be included in the laboratory data package submitted to the TCEQ in the TRRP-required report(s).

Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.

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Appe	ndix A (cont'd): Laboratory Review	Checklist: Exception Reports
Laborat	tory Name: STL-Houston	LRC Date: 06/26/06
Project Name: 9003 N. MAIN		Laboratory Job Number: 318054
Reviewer Name: ERA		Prep Batch Number(s): 157626 (Water)-BTEX
ER# ¹	DESCRIPTION	
1	The temperature of the cooler received b	y the laboratory on 06/21/06 was above the acceptable range of 2.0-6.0 °C.
2		veries on both columns in sample 318054-4 were above acceptance limits
3	The laboratory selected another client's s	ample to perform as the MS/MSD.
4	The benzene RPD between the two colur lower of the two results was reported.	nns in sample 318054-2 was >40%. Since anomalies were present, the

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked on the LRC)

Lab	orato	ry Name; STL-Houston LRC	C Date: 06/26/06					
Proi	ect N	lame: 9003 N. MAIN Lab	oratory Job Number: 318054	••••••				
,				nton) E	·	aldah	do	
#1	A ²		Batch Number(s): 157516 (soil and W					ED 45
#	A ^s	Description12		Yes	No	NA.	NK	EK#
		Chain-of-custody (C-O-C)						
R1	OI	Did samples meet the laboratory's standard conditions of samp			<u>X</u>	ļ		1
		Were all departures from standard conditions described in an ex-	xception report?	X		<u> </u>		
R2	OI	Sample and quality control (QC) identification		1		L		
		Are all field sample ID numbers cross-referenced to the laborat		X		ļ		
	- T	Are all laboratory ID numbers cross-referenced to the correspo	nding QC data?	X		ļ	A3 NR4	ļ
R3	OI	Test reports		1		ļ	ļ	ļ
		Were all samples prepared and analyzed within holding times?		X				
		Other than those results < MQL, were all other raw values brace	keted by calibration standards?	X		 -		
		Were calculations checked by a peer or supervisor? Were all analyte identifications checked by a peer or superviso	w?	$\frac{\Lambda}{X}$				
		Were an analyte identifications checked by a peer or superviso. Were sample quantitation limits reported for all analytes not de		$\frac{X}{X}$				
		Were all results for soil and sediment samples reported on a dry		X				
		Were % moisture (or solids) reported for all soil and sediment s		$\frac{\hat{x}}{x}$				
		If required for the project, TICs reported?	samprest	+^+		X		
R4	0	Surrogate recovery data		++		<u>^`</u>	A ³ NR ⁴	
	<u> </u>	Were surrogates added prior to extraction?		$ \mathbf{x} $				·
		Were surrogate percent recoveries in all samples within the lab	oratory OC limits?	$\frac{1}{x}$				
R5	OI	Test reports/summary forms for blank samples		1 1				
		Were appropriate type(s) of blanks analyzed?		X				
		Were blanks analyzed at the appropriate frequency?		X				
		Were method blanks taken through the entire analytical process	s, including preparation and, if	X				
		applicable, cleanup procedures?						
		Were blank concentrations < MQL?		X				
R6	OI	Laboratory control samples (LCS):						
		Were all COCs included in the LCS?		X				
		Was each LCS taken through the entire analytical procedure, in	cluding prep and cleanup steps?	X				
		Were LCSs analyzed at the required frequency?		X				
		Were LCS (and LCSD, if applicable) %Rs within the laborator		X				***********
		Does the detectability data document the laboratory's capability	y to detect the COCs at the MDL used			X		
		to calculate the SQLs?		↓				
	<u> </u>	Was the LCSD RPD within QC limits?		11		X		
R7	OI	Matrix spike (MS) and matrix spike duplicate (MSD) data	1 Lang	 ,, 				
		Were the project/method specified analytes included in the MS	and MSD?	X			XXX	
		Were MS/MSD analyzed at the appropriate frequency? Were MS (and MSD, if applicable) %Rs within the laboratory of	OC 1::	X			V	
			QC limits?	+-+				2
R8	OI	Were MS/MSD RPDs within laboratory QC limits? Analytical duplicate data		+-+			 	
10	01	Were appropriate analytical duplicates analyzed for each matrix	······································	╂╼╂		х		
		Were analytical duplicates analyzed at the appropriate frequence		╂──╂		X		
		Were RPDs or relative standard deviations within the laboratory		╁╌┼		$\frac{\Lambda}{X}$		
R9	OI	Method quantitation limits (MQLs):	y QC imms:	\vdash		^		
		Are the MQLs for each method analyte included in the laborato	ry data package?	$ \mathbf{x} $				
		Do the MQLs correspond to the concentration of the lowest nor		$\frac{x}{x}$		-		
1		Are unadjusted MQLs included in the laboratory data package?		$\frac{1}{X}$	\neg	$\neg \uparrow$	$\neg \neg$	
R10		Other problems/anomalies		T		一	$\neg \dagger$	
		Are all known problems/anomalies/special conditions noted in t	this LRC and ER?	x			$\neg \uparrow$	
		Were all necessary corrective actions performed for the reported		X				
		Was applicable and available technology used to lower the SQI	to minimize the matrix interference	X	\neg			

^{1.} Items identified by the letter "R" must be included in the laboratory data package submitted in the TRRP-required report(s). Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.

ΑI

⁼ organic analyses; I = inorganic analyses (and general chemistry, when applicable);

NA = Not applicable;
NR = Not reviewed;
ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

	-	ndix A (cont'd): Laboratory Review Checklist: Reportable Data	<u>a</u>					
		ory Name: STL-Houston LRC Date: 06/26/06						
Proj	ect N	Name: 9003 N. MAIN Laboratory Job Number: 31	8054					
Rev	iewe	Prep Batch Number(s): 157:	516 (soil and V	Vate	r)-Fo	rmald	ehyde	
#1	A ²	Description		Yes		NA ³	NR ⁴	ER#
		Initial calibration (ICAL)				15/11/2	7 6 37	10.00
<u> </u>	<u>~-</u>	Were response factors and/or relative response factors for each analyte within OC limits?	,	X		 	 	1
		Were percent RSDs or correlation coefficient criteria met?		X				
		Was the number of standards recommended in the method used for all analytes?		X		 		†
		Were all points generated between the lowest and highest standard used to calculate the c	urve?	X				1
		Are ICAL data available for all instruments used?		X			<u> </u>	1
		Has the initial calibration curve been verified using an appropriate second source standar	d?	X			 	1
S2	OI	Initial and continuing calibration verification (ICCV and CCV) and continuing calil			11111	1 1 1 1		1 3 4
		Was the CCV analyzed at the method-required frequency?		X				1
		Were percent differences for each analyte within the method-required QC limits?		X			1	1
		Was the ICAL curve verified for each analyte?		X		<u> </u>	T	1
		Was the absolute value of the analyte concentration in the inorganic CCB < MDL?				X		
S3	0	Mass spectral tuning:	· · · · · · · · · · · · · · · · · · ·				11.1	1
		Was the appropriate compound for the method used for tuning?				X	†	1
		Were ion abundance data within the method-required QC limits?				X	1	1
S4	0	Internal standards (IS):	····	1.5-1.5	1, 1,	2 (3)		
-		Were IS area counts and retention times within the method-required QC limits?				X		
S5	OI	Raw data (NELAC section 1 appendix A glossary, and section 5.12 or ISO/IEC 1702	5 section	1.				1
	<u> </u>	Were the raw data (for example, chromatograms, spectral data) reviewed by an analyst?		X			1	1
		Were data associated with manual integrations flagged on the raw data?		X			1	
S6	0	Dual column confirmation						
		Did dual column confirmation results meet the method-required QC?				X	1	1
S7	0	Tentatively identified compounds (TICs):	<u> </u>				1	1
		If TICs were requested, were the mass spectra and TIC data subject to appropriate checks	?			X	1	
S8	7	Interference Check Sample (ICS) results:		77.77			 	1
		Were percent recoveries within method QC limits?				X		1
S9	Ī	Serial dilutions, post digestion spikes, and method of standard additions		5,117	33.4	2.85	100	1
 -		Were percent differences, recoveries, and the linearity within the QC limits specified in t	he method?			X		1
S10	OI	Method detection limit (MDL) studies		100	100	200	155.75	
	~-	Was a MDL study performed for each reported analyte?		X				1
		Is the MDL either adjusted or supported by the analysis of DCSs?		X				1
S11	OI	Proficiency test reports:		120	720.0	Alter.	44.5	VA,
		Was the laboratory's performance acceptable on the applicable proficiency tests or evaluation	tion studies?	X				1
S12	OI	Standards documentation		1.7			11.57	
		Are all standards used in the analyses NIST-traceable or obtained from other appropriate	sources?	X				T
S13	OI	Compound/analyte identification procedures						
		Are the procedures for compound/analyte identification documented?		X				1
S14	OI	Demonstration of analyst competency (DOC)						T
		Was DOC conducted consistent with NELAC Chapter 5C or ISO/IEC 4?		X				
İ		Is documentation of the analyst's competency up-to-date and on file?		X				
S15	OI	Verification/validation documentation for methods (NELAC Chap 5 or ISO/IEC 1702	5 Section 5)		-7,71		1.57 11.	1
		Are all the methods used to generate the data documented, verified, and validated, where		X				Τ
~	OI	Laboratory standard operating procedures (SOPs):			. 1741	77.7	F 1 1 2 2	
SIM	1 31							

Items identified by the letter "R" should be included in the laboratory data package submitted to the TCEQ in the TRRP-required report(s). Items identified by the letter "S" should be retained and made available upon request for the appropriate retention period.

O = organic analyses; I = inorganic analyses (and general chemistry, when applicable).

NA = Not applicable.

NR = Not Reviewed.

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked).

Apper	ndix A (cont'd): Laboratory Revie	ew Checklist: Exception Reports
Laborat	ory Name: STL-Houston	LRC Date: 06/26/06
Project Name: 9003 N. MAIN Laboratory Job Number: 318054		
Review	er Name: JPS	Prep Batch Number(s): 157516 (soil and Water)-Formaldehyde
ER#1	DESCRIPTION	
1	The temperature of the cooler received	d by the laboratory on 06/21/06 was above the acceptable range of 2.0-6.0 °C.
2	The laboratory selected another client	's sample to perform as the MS/MSD.

ER# = Exception Report identification number (an Exception Report should be completed for an item if "NR" or "No" is checked on the LRC)

SEVERN STL

CHAIN OF CUSTODY RECORD

Gustomet Information	Q.			; ;		6
1	PROJECT NAME	31-5.6.10.11	9003 N W .	76.0 76.0		,
0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0		IE	'V', 'V'\a'W	O C: 8315 F01	B.TPH TX 1005 C:8315 Formaldehyde	
COMPANY ATC Associates, inc.	BILL TO	ATC Associates, Inc.	,		,	
SEND REPORT TO Patrick Duoraczyk	INVOICE ATTIN	Patrick Dworaczyk			solis bry meighter	P karina kalauna kar
AODRESS 3928 Blaebonnet Drive	ADDRESS 39	3928 Bluedonnet Orive		т		
				- × -		
				J Z Z	0,8054	
GITWSTATEZIE Stafford, Il 77477	CITY/STATEZIP	Stafford, TX 77477		2 O a	7 0 1	************
PHONE: 281-240-0154	PHONE 28	281-240-0154		701		
FAX 281-240-8909	FAX. 8.28.	281-240-8909		rω		***************************************
SAMP NO SAMPLE DESCRIPTION	PRESERVE	F SAMPLE MATRIX SA	SAMPLE DATE SA	SAMPLE TIME # CONTAINER	ER A B C D E F G H I J K L M NO	POR
3-1 12-14		7.98	90-12-9	950		
B-2 20-22			╂	1100		
B3 20-22		Sp;		11.55		
<u>B.1</u>		Water		arci	7,7,7	
[የጋ		Water	<u> </u>	1245	N.V.	
3-3		Water		001		
Sampler:	Shipment Method: Ares of	Airbill	No.:		Required TurnAround: 5	
1. Relinquished By Reference 1	Date 2. Relinquished By:		Date	3. Relinquished By:		Date
Company Name:	Time Company Name:		Time	Company Name:	<u></u>	Time
1. Received By: GMTD	pate 2. Received By:		Date	3. Received By:	0	Date
Company Name:	Time, 24 Company Name:		Time	Company Name:		Time
19			1			

713-690-4444 RAX 713-690-5646 Bouston, TX 77040 Severn Trent Laboratories 6310 Rothway Brive

rpjsckl	Job Sample Receipt Checklist Report	V2
Customer Job ID:	n.: 57216 Check List Number.: 1 Description.: Job Check List Date.: 06/21/2006 ject Description.: TRRP Project , Inc. Contact.: Patrick Dwora	Date of the Report: 06/21/2006 Project Manager: daj1 czyk
Questions ?	(Y/N) Comments	
Chain of Custody Received?	Y	
If "yes", completed properly	? Y	
Custody seal on shipping contain	ner? N	
If "yes", custody seal intac	t?	
Custody seals on sample contain	ers? N	
If "yes", custody seal intac	t?	
Samples chilled?	N see src	
Temperature of cooler acceptabl	e? (4 deg C +/- 2). N 18.6	
If "no", is sample an air ma	trix?(no temp req.) N	
Thermometer ID	Y 437	2 /
Samples received intact (good c	ondition)? Y	()(
Volatile samples acceptable? (n	o headspace) Y	10-21-6
Correct containers used?	Y	10-1/1
Adequate sample volume provided	17Y	V
Samples preserved correctly?	Y	
Samples received within holding	y-time? Y y	
Agreement between COC and sampl	e labels? Y	
Radioactivity at or below backg	round levels? Y	
Additional		
Comments		
Sample Custodian Signature/Date	Y jac	

Page 1

V	4 Tre			- SAMPLE		CHECKLIS	r/, ,
CLIENT NAME:	/ / (<u> </u>	<u> </u>		CARRIE	ER/DRIVER	NAME:	Client
PROJECT:	·	·	UN	PACKED BY:	Profession		
DATE RECEIVED:			***************************************		UNPACKE	ED STAMP:	
TOTAL # COOLER	S RECEIVED):					
<u> </u>				OLER CHE	CKLIST		
COOLER ID	COC PRESENT	CUSTOD	Y TAPE	COOLER TEMP	THERM ID	TEMP BLK PRESENT	List Sample Bottles in Each Cooler if out of Temperature
	(Y/N)	PRESENT (Y/N)	INTACT (Y/N)	(°C)	טו	(Y/N)	out of remperature
Gray lihit,	y	C B/V	N	18.6	437	N	Chillippograss
	,	С	· · · · · · · · · · · · · · · · · · ·		·		11/5
		В					
		С					
C=COOLER B	= BOTTLES	В					
** SPECIFIC PROJEC VOLATILE HEADSF (If ANY headspace is p pH OF WATER SAM PRESERV H2SO4 (<2) HNO3 (<2) HCL (<2) (Not VOA NaOH – Cyanide (> NaOH/Zn Acetate – Other	PACE ACCEF resent, list det MPLES /ATION Vials)	PTABLE? Yes ails in INCONSI	No	NA	Je M N	umber of VO	**************************************
# OF NEAT BOTTLE	ES:				#	OF SOIL JAF	RS:
		INCONSIST	ENCIES (Place in Job	Notes as v	vell (CTRL F	-12)
PERSON CONTACT RESOLUTION	ED:			ACTION TAP		DATE:	
NOTES							
Project Manager					****	(Use b	ack of sheet if necessary)



7/10/2006 Date:

> B-1 12-14 Customer Sample ID:

Job Number: 318054

Date/Time Sampled 6/21/2006

09:50 13:31 Date/Time Received: 6/21/2006

Laboratory Sample ID: 318054-001

Sample Matrix Soil

			Value	S WIDT	MOIN	108		Analysis Date June	Batch	10	Vralyst
Method: SM-2540 G Mod., Soil											
% Solids	NA	82.2				aller en Production en	%	6/22/2006 16:30	157440	1.00	ydh
Moisture	MOIST	17.8				and the second second	%	6/22/2006 16:30	157440	1.00	yps
Method: SW-846 8021B, Soil											
Benzene	71-43-2	5.60	D D	4.60	10.0	5.60	ug/Kg	6/23/2006 20:50	157581	1.00	era
Ethylbenzene	100-41-4	7.42	Þ	6.10	10.0	7.42	ug/Kg	6/23/2006 20:50	157581	1.00	era
m,p-Xylene	MPXYLENE	15.4	Þ	12.7	20.0	15.4	ug/Kg	6/23/2006 20:50	157581	1.00	era
o-Xylene	95-47-6	7.86	Ð	6.46	10.0	7.86	ug/Kg	6/23/2006 20:50	157581	1.00	ега
Toluene	108-88-3	7.74	D	6.36	10.0	7.74	ug/Kg	6/23/2006 20:50	157581	1.00	era
Xylenes (total)	1330-20-7	23.3	D	19.1	30.0	23.3	ug/Kg	6/23/2006 20:50	157581	1.00	era
Method: SW-846 8315, Soil											
Formaldehyde	0-00-09	2520		2.87	100	6.69	ug/Kg	6/26/2006 11:56	157630	1.00	jps
Method: SW846 8315, Water											
Solid Phase Extraction	NA	Complete					N/A	6/23/2006 08:00	157516	1.00	oue
									•		***************************************

Form I



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-1 12-14

Date/Time Sampled: 6/21/2006 09:50

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-001

Sample Matrix Soil

D. E. H. Analysti		1.00 mep	I.00 mep	1.00 mep	1.00 mep		
Batch		157465	157465	157465	157465		157296
AnalysisDareTime		6/22/2006 20:38	6/22/2006 20:38	6/22/2006 20:38	6/22/2006 20:38		6/21/2006 16:00
SUND			mg/Kg	mg/Kg	mg/Kg		N/A
NOS		13.8	13.8	6.57	13.8		
MOI		50.0	50.0	20.0	20.0		
MDI		11.4	11.4	5.40	11.4		
		þ	n	D	n		
NESC IN		13.8	13.8	6.57	13.8		Complete
		NA	NA	NA	NA		N A
	TNRCC 1005, Soil	Petroleum Hydrocarbons C12 - C28	Petroleum Hydrocarbons C28 - C35	Petroleum Hydrocarbons C6 - C12		Method: TNRCC TX-1005, Soil	TNRCC 1005 Extraction

Form I



Date: 7/10/2006

Customer Sample ID: B-2 20-22

Job Number: 318054

Date/Time Sampled 6/21/2006 11:00

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-002

Sample Matrix Soil

	Svo	NESS COL		MDE		108		Analysis Date mile	Batch	DE	Analyst
Method: SM-2540 G Mod., Soil											
% Solids	NA	85.2			-	:	%	6/22/2006 16:30	157440	1.00	sdħ
Moisture	MOIST	14.8					%	6/22/2006 16:30	157440	1.00	sdħ
Method: SW-846 8021B, Soil											
Benzene	71-43-2	72.0		4.60	10.0	5.40	ug/Kg	6/23/2006 21:30	157581	1.00	era
Ethylbenzene	100-41-4	785		6.10	10.0	7.16	ug/Kg	6/23/2006 21:30	157581	1.00	era
m,p-Xylene	MPXYLENE	1450		12.7	20.0	14,9	ug/Kg	6/23/2006 21:30	157581	1.00	era
o-Xylene	95-47-6	512		6.46	10.0	7.58	ug/Kg	6/23/2006 21:30	157581	1.00	era
Toluene	108-88-3	155		6.36	10.0	7.46	ug/Kg	6/23/2006 21:30	157581	1.00	era
Xylenes (total)	1330-20-7	1970		19.1	30.0	22.5	ug/Kg	6/23/2006 21:30	157581	1.00	era
Method: SW-846 8315, Soil											
Formaldehyde	20-00-0	575		2.87	100	67.3	ug/Kg	6/26/2006 12:12	157630	1.00	jps
Method: SW846 8315, Water											
Solid Phase Extraction	NA	Complete					N/A	6/23/2006 08:00	157516	1.00	enc

Form I



Date: 7/10/2006

Customer Sample ID: B-2 20-22

Job Number: 318054

Date/Time Sampled 6/21/2006 11:00

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-002

Sample Matrix Soil

COPER S						 		
analyst.		mep	mep	mep	mep		lvp	
		1.00	1.00	1.00	1.00		1.00	
Batch		157465	157465	157465	157465		157296	
Dole Inc		6/22/2006 21:44	6/22/2006 21:44	6/22/2006 21:44	6/22/2006 21:44		6/21/2006 16:00	
Affilia		6/22/	6/22/	6/22/	6/22/		6/21/	
A VIOLET SOLUTION DE LA MINE AMBESTS DATE MANAGEMENT		mg/Kg	mg/Kg	mg/Kg	mg/Kg		N/A	
SOI		13.3	13.3	6.32	13.3			
NO		20.0	50.0	50.0	50.0			
MDL		11.4	11.4	5.40	11.4			
Ø.		<u> </u>	<u> </u>	<u> </u>	Þ			
RESULT OF		13.3	13.3	6.32	13.3		Complete	
English Substitute		NA	NA	NA	NA		NA V	
	Method: TNRCC 1005, Soil	Petroleum Hydrocarbons C12 - C28	Petroleum Hydrocarbons C28 - C35	Petroleum Hydrocarbons C6 - C12	Petroleum Hydrocarbons C6 - C35	Method: TNRCC TX-1005, Soil	TNRCC 1005 Extraction	
	Meth	Petrole	Petrole	Petrol	Petrole	Meth	TINEC	

Form I



Date: 7/10/2006

USECMEN PARCEASSOCIAL E

Job Number: 318054

Customer Sample ID: B-3 20-22

Date/Time Sampled: 6/21/2006 11:55

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-003

Sample Matrix Soil

	To AS T	RESULT		JUN	MOLE	SOL		Makis Bate mine	Batch		Analysi
Method: SM-2540 G Mod., Soil											
% Solids	NA	85.3					%	6/22/2006 16:30	157440	1.00	sdh
Moisture	MOIST	14.7		to reconstruction			%	6/22/2006 16:30	157440	1.00	ydh
Method: SW-846 8021B, Soil											
Benzene	71-43-2	5.39	5	4.60	10.0	5.39	ug/Kg	6/23/2006 22:30	157581	1.00	era
Ethylbenzene	100-41-4	7.15	D D	6.10	10.0	7.15	ug/Kg	6/23/2006 22:30	157581	1.00	era
m,p-Xylene	MPXYLENE	14.9		12.7	20.0	14.9	ug/Kg	6/23/2006 22:30	157581	1.00	era
o-Xylene	95-47-6	7.57		6.46	10.0	7.57	ug/Kg	6/23/2006 22:30	157581	1.00	era
Toluene	108-88-3	7.45	D	6.36	10.0	7.45	ug/Kg	6/23/2006 22:30	157581	1.00	era
Xylenes (total)	1330-20-7	22.4	n	19.1	30.0	22.4	ug/Kg	6/23/2006 22:30	157581	1.00	era
Method: SW-846 8315, Soil											
Formaldehyde	20-00-0	357		2.87	100	67.3	ug/Kg	6/26/2006 12:27	157630	1.00	sdí
Method: SW846 8315, Water											
Solid Phase Extraction	NA	Complete					N/A	6/23/2006 08:00	157516	1.00	enc

Form I



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-3 20-22

Date/Time Sampled: 6/21/2006

11:55

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-003

Sample Matrix Soil

Set State		0 mep	0 mep	0 mep	0 mep		0 Ivp	
G		1.00	1.00	1.00	1.00		1.00	
Batch		157465	157465	157465	157465		157296	
Analysis Date Illines Barel		6/22/2006 22:17	6/22/2006 22:17	6/22/2006 22:17	6/22/2006 22:17		6/21/2006 16:00	
Analysis		6/22/20	6/22/20	6/22/20	6/22/20		6/21/20	
		mg/Kg	mg/Kg	mg/Kg	mg/Kg		N/A	
TOS		13.3	13.3	6.33	13.3			
TON!		50.0	50.0	50.0	50.0			
VO		11.4	11.4	5.40	11.4			
5)								·
0		Ω	Þ		ם		<u></u>	
KESU		13.3	13.3	6.33	13.3		Complete	
		¥	4	∢	₹		¥	
		NA	NA	N A	NA		NA	
		C28	C35	212	335	Soil		
TO STOWER THOUSA	Method: TNRCC 1005, Soil	Petroleum Hydrocarbons C12 - C28	Petroleum Hydrocarbons C28 - C35	Petroleum Hydrocarbons C6 - C12	Petroleum Hydrocarbons C6 - C35	Method: TNRCC TX-1005, Soil	ď	
	3C 186	carbon	carbon	carbon	carbon:	XI X	TNRCC 1005 Extraction	
	TNRC	Hydroc	Hydro	Hydroc	Hydroc	TNRC	05 Exi	
	iod:	leum l	leum l	leum]	leum J	; poq:	CC 10	
	Meti	Petro	Petro	Petro	Petro	Met	TAR	

Form I



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-1

Date/Time Sampled: 6/21/2006 12:10

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-004

Sample Matrix Water

THE STANFORD STANFORD		RESULT	9 FE A.G		MOL	Sol		Analysis Dale Time	Barch	. TO	Malya
Method: SW-846 8021B, Water											
Benzene	71-43-2	25.6	·	0.170	0.500	0.170	ng/L	6/26/2006 10:20	157626	1.00	era
Ethylbenzene	100-41-4	131		0.230	0.500	0.230	T/gn	6/26/2006 10:20	157626	1.00	era
m,p-Xylene	MPXYLENE	297		0.370	1.00	0.370	ng/L	6/26/2006 10:20	157626	1.00	era
o-Xylene	95-47-6	6.66		0.190	0.500	0.190	T/Sn	6/26/2006 10:20	157626	1.00	ета
Toluene	108-88-3	7.05		0.190	0.500	0.190	ng/L	6/26/2006 10:20	157626	1.00	ета
Method: SW846 8315, Water											
Solid Phase Extraction	NA	Complete					N/A	6/23/2006 08:00	157516	1.00	enc
Method: SW-846 8315, Water											
Formaldehyde	20-00-0	27.6		2.87	5.00	3.38	T/fin	6/26/2006 12:58	157630	1.00	sdí
Method: TNRCC 1005, Water											
Petroleum Hydrocarbons C12 - C28	NA	0.830		0.870	2.00	0.830	mg/L	6/23/2006 23:03	157591	1.00	mep
Petroleum Hydrocarbons C28 - C35	NA	0.830		0.870	5.00	0.830	mg/L	6/23/2006 23:03	157591	1.00	шер
Petroleum Hydrocarbons C6 - C12	NA	0.480	ם	0.500	2.00	0.480	mg/L	6/23/2006 23:03	157591	1.00	mep
Petroleum Hydrocarbons C6 - C35	NA	0.830	D	0.870	5.00	0.830	mg/L	6/23/2006 23:03	157591	1.00	dəm
-											

Form I

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Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-1

Date/Time Sampled 6/21/2006 12:10

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-004

Sample Matrix Water

Method: TNRCCTX-1005, Water	TNRCC 1005 Extraction
	ž
	Complete
	A/N
	6/22/2006 14:30
	157449
	1.00 lvp

Form I



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-2.

Date/Time Sampled: 6/21/2006 12:45

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-005

Sample Matrix Water

THE WILLIAM DESCRIPTION OF THE PROPERTY OF THE		THE SERVICE OF		KOJA JA	IOM	NOS.	SHAULE	Analysis Date Trine	Batch	DIC	A CHELLY ST
Method: SW-846 8021B, Water											
Benzene	71-43-2	0.170	n	0.170	0.500	0.170	ng/L	6/26/2006 10:40	157626	1.00	era
Ethylbenzene	100-41-4	0.670		0.230	0.500	0.230	ng/L	6/26/2006 10:40	157626	1.00	era
m,p-Xylene	MPXYLENE	1.81		0.370	1.00	0.370	ng/L	6/26/2006 10:40	157626	1.00	era
o-Xylene	95-47-6	0.560	***************************************	0.190	0.500	0.190	J/gn	6/26/2006 10:40	157626	1.00	era
Toluene	108-88-3	0.190	n	0.190	0.500	0.190	ug/L	6/26/2006 10:40	157626	1.00	era
Method: SW846 8315, Water											
Solid Phase Extraction	NA	Complete					N/A	6/23/2006 08:00	157516	1.00	enc
Method: SW-846 8315, Water											
Formaldehyde	20-00-0	46.4		2.87	5.00	3.38	ng/L	6/26/2006 13:13	157630	1.00	ips
Method: TNRCC 1005, Water											
Petroleum Hydrocarbons C12 - C28	NA	0.830	n	0.870	5.00	0.830	mg/L	6/23/2006 23:36	157591	1.00	mep
Petroleum Hydrocarbons C28 - C35	NA	0.830	n	0.870	2.00	0.830	mg/L	6/23/2006 23:36	157591	1.00	mep
Petroleum Hydrocarbons C6 - C12	NA	0.480	D	0.500	5.00	0.480	mg/L	6/23/2006 23:36	157591	1.00	mep
Petroleum Hydrocarbons C6 - C35	NA	0.830	ב	0.870	5.00	0.830	mg/L	6/23/2006 23:36	157591	1.00	mep
		****		•	-	-					_

Form I



7/10/2006 Date:

Job Number: 318054

Customer Sample ID:

12:45 Date/Time Sampled 6/21/2006

13:31 Date/Time Received 6/21/2006

Laboratory Sample ID: 318054-005

Sample Matrix Water

Contraction de la contraction		
		qv
300		1.00
Bates.		157449
THIC		
Spirit and		6/22/2006 14:30
		6/22
		N/A
		-
108		
Military		
)		
		Сотрієє
		СО
188		
		₹ Z
	Vater	
0.01	-1005, V	
	CCTX	TNRCC 1005 Extraction
	t: TNR	1005 E
	Methoc	TNRCC
	1 T T	

Form I



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-3

Date/Time Sampled 6/21/2006 13:00

Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-006

Sample Matrix Water

HAND TOURS IN SERVICE		KASIII T		GIV NO	MOI	10S	TWIS 4	Analysis Date turie		110	finalyst
Method: SW-846 8021B, Water											
Benzene	71-43-2	0.170	<u></u>	0.170	0.500	0.170	ng/L	6/26/2006 11:57	157626	1.00	era
Ethylbenzene	100-41-4	0.230	D	0.230	0.500	0.230	ng/L	6/26/2006 11:57	157626	1.00	era
m,p-Xylene	MPXYLENE	0.370	D	0.370	1.00	0.370	ng/L	6/26/2006 11:57	157626	1.00	ета
o-Xylene	95-47-6	0.190		0.190	0.500	0.190	T/Sn	6/26/2006 11:57	157626	1.00	cra
Toluene	108-88-3	0.190	D	0.190	0.500	0.190	ug/L	6/26/2006 11:57	157626	1.00	era
Method: SW846 8315, Water											
Solid Phase Extraction	NA	Complete		tan anka k			N/A	6/23/2006 08:00	157516	1.00	enc
Method: SW-846 8315, Water											
Formaldehyde	20-00-0	9.80		2.87	5.00	3.54	ng/L	6/26/2006 13:29	157630	1.00	jps
Method: TNRCC 1005, Water											
Petroleum Hydrocarbons C12 - C28	NA	0.840	n n	0.870	2.00	0.840	mg/L	6/24/2006 00:09	157591	1.00	шер
Petroleum Hydrocarbons C28 - C35	NA	0.840	Þ	0.870	5.00	0.840	mg/L	6/24/2006 00:09	157591	1.00	mcp
Petroleum Hydrocarbons C6 - C12	NA	0.480	n	0.500	5.00	0.480	mg/L	6/24/2006 00:09	157591	1.00	mep
Petroleum Hydrocarbons C6 - C35	NA	0.840)	0.870	5.00	0.840	mg/L	6/24/2006 00:09	157591	1.00	mep

Form I



Date: 7/10/2006

Job Number: 318054

Customer Sample ID: B-3

Date/Time Sampled 6/21/2006 13:00

.Date/Time Received: 6/21/2006 13:31

Laboratory Sample ID: 318054-006

Sample Matrix Water

lvp

1.00

157449

6/22/2006 14:30

N/A

Complete NA Method: TNRCC TX-1005, Water TNRCC 1005 Extraction

Form 1



QUALITY CONTROL RESULTS

Job Number.: 318054

Report Date.: 07/10/2006

CUSTOMER: ATC Associates, Inc.

PROJECT: 9003 N. MAIN

ATTN: Patrick Dworaczyk

Mε	est Method thod Descrip arameter	otion,: Moi	sture (Total +	Fixed Solids,					st: sdh Code.: %SOLID	
QC	Lab ID	Reagent	QC Result	QC Result	True Value	Orig. Value	Calc. Result *	Limits	F Date	Tim
DU	318054-3		85.8368	<u> </u>		85.3351	0.6	10.0	06/22/2006	163
UC	318174-6		83.8722			83.3475	0.6	10.0	06/22/2006	163
U	318107-4		98.3661			98.3314	0.0	10.0	06/22/2006	163
DU	318174-16		85.3161			85.0272	0.3	10.0	06/22/2006	163
MB	15744021		0.0000						06/22/2006	163
MB	15744021		0.0000						06/22/2006	163
MB	15744021		0.0000						06/22/2006	
DU	318100-8		79.4863			78.5088	1.2	10.0	06/22/2006	163
DU	318146- 1		15.6110			15.7773	1.1	10.0	06/22/2006	163
Me Pe	irameter	otion.: Moi	sture (Total + sture	Fixed Solids,	Batch(s): 157440		Test	st: sdh Code.: MOIST	7:-
SC	Lab ID	Reagent	QC Result	QC Result	True Value	Orig. Value	Calc. Result *	Limits	F Date	Tin
U	318174-16		14.6839			14.9728	1.9	10.0	06/22/2006	
U	318174-6		16.1278			16.6525	3.2	10.0	06/22/2006	
UC	318100-8		20.5137			21.4912	4.7	10.0	06/22/2006	
U	318107-4		1.6339			1.6686	2.1	10.0	06/22/2006	
UC	318054-3		14.1632			14.6649	3.5	10.0	06/22/2006	
บเ	318146-1		84.3890			84.2227	0.2	10.0	06/22/2006	4/"



m,p-Xylene, Soil
o-Xylene, Soil
Xylenes (total), Soil
Total BTEX, Soil
Tert-Butyl Methyl Ether Column B, Soil

Methyl tert-Butyl ether, Soil

Benzene, Soil Toluene, Soil Ethylbenzene, Soil

Benzene Column B, Soil

Job Number.: 318054	QUALITY	CONTRO	. RESULI		eport Date.: 07	7/10/2006	
CUSTOMER: ATC Associates, Inc.	PROJE	CT: 9003 N. MA	i N	A	TTN: Patrick Dv	ioraczyk	
QC Type Description		Reag. Cod	e Lab	ID	Dilution Factor	Date	Time
Test Method: SW-846 80218 Method Description.: GC Volatile Organi	cs		: uç): 157581 1		Anal	yst: era	
LCS Laboratory Control Sample		BX\$0612060	157581-	ı		06/23/200	1709
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Va	lue Calc. Res	sult * Limi	ts F
Methyl tert-Butyl ether, Soil Benzene, Soil Toluene, Soil Ethylbenzene, Soil m,p-Xylene, Soil o-Xylene, Soil Xylenes (total), Soil Total BTEX, Soil Tert-Butyl Methyl Ether Column B, Soil Benzene Column B, Soil Toluene Column B, Soil Ethylbenzene Column B, Soil ethylbenzene Column B, Soil o-Xylene Column B, Soil	46.6816 52.0285 52.1341 52.0014 109.893 55.4779 166.0887 322.2527 45.1083 49.6298 51.3092 51.3521 104.727 56.1957		50.00000 50.00000 50.00000 100.00000 150.00000 150.00000 50.00000 50.00000 50.00000 50.00000 50.00000		93.4 104.1 104.3 109.9 111.0 110.7 107.4 90.2 99.3 102.6 102.7 104.7	69- 70- 71- 70- 70- 70- 61- 69- 71- 71-	125 133 134 139 136 131 130 130 125 133 134 139 136
MB Method Blank			157581-	1		06/23/200	6 1730
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Va	ilue Calc. Re	sult * Limi	its F
Methyl tert-Butyl ether, Soil Benzene, Soil Toluene, Soil Ethylbenzene, Soil m,p-Xylene, Soil o-Xylene, Soil Yotal BTEX, Soil Tert-Butyl Methyl Ether Column B, Soil Benzene Column B, Soil Toluene Column B, Soil Ethylbenzene Column B, Soil m,p-Xylene Column B, Soil o-Xylene Column B, Soil	ND ND ND ND ND 0.0000 ND ND ND ND ND ND						
SB Spiked Blank		BXS061206C	157581+	1		06/23/200	06 1930
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Va	olue Calc. Re	sult * Lim	its F

Page 35	*	%=% REC.	R=RPD, A=AB	S Diff.,	D=% Diff.

ND

ND

ND

ND

ND

ND

ND

ND

0.0000

0.0000

93

101

102

100

105

106

110

107

91

94

50.000000

50.000000

50.000000

50.000000

100.000000

50.000000

150.000000 300.000000

50.000000

50.000000

30.0-130.0

30.0-130.0 30.0-130.0 30.0-130.0

30.0-130.0

30.0-130.0

30.0-130.0

30.0-130.0

30.0-130.0

30.0-130.0

46.6336

50.6572 51.1370

50.0466

53.1742

165.5110

320.0681

45.5755

47.0232

104.969



Job Number.: 318054	QUALITY	CONTROL	RESULT		Date.: 07/10)/2006	
CUSTOMER: ATC Associates, Inc.	PROJEC	T: 9003 N. MAIN		ATTN:			
QC Type Description		Reag. Code	Lab	ID Dilut	ion Factor	Date Time	<u>.</u>
SB Spiked:Blank		BXS061206C	157581-1			06/23/2006 193	50
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Toluene Column B, Soil Ethylbenzene Column B, Soil m,p-Xylene Column B, Soil o-Xylene Column B, Soil	48.2890 52.7629 100.260 60.5420		50.000000 50.000000 100.000000 50.000000	ND ND ND ND	97 106 100 121	30.0-130.0 30.0-130.0 30.0-130.0 30.0-130.0	
SBD Spiked Blank Duplicate		BX\$061206C	157581-1			06/23/2006 195	. 0
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	: * Limits	F
Methyl tert-Butyl ether, Soil	49.6281	46.6336	50.000000	ND	99.3 6.2	30-130 20	
Benzene, Soil	54.0149	50.6572	50.000000	ND	108.0 6.4	30-130 20	
Toluene, Soil	54.0783	51.1370	50.000000	ND	108.2 5.6	30-130 20	
Ethylbenzene, Soil	54.2439	50.0466	50.000000	ND	108.5 8.0	30-130 20	
m,p-Xylene, Soil	113.256	104.969	100.000000	ND	113.3	30-130 20	
o-Xylene, Soil	56.5970	53.1742	50.000000	ND	7.6 113.2	30-130	
Xylenes (total), Soil	169.8530	165.5110	150.000000	0.0000	6.2 113.2	20 30-130	
Total BTEX, Soil	332.1901	320.0681	300.000000	0.0000	2.6 110.7	20 30-130	
Tert-Butyl Methyl Ether Column B, Soil	48.6404	45.5755	50.000000	ND	3.7 97.3	20 30-130	
Benzene Column B, Soil	50.4222	47.0232	50.000000	ND	6.5 100.8	20 30-130	
Toluene Column B, Soil	51.9796	48.2890	50.000000	ND	7.0 104.0	20 30-130	
Ethylbenzene Column B, Soil	53.2134	52.7629	50.000000	ND	7.4 106.4	20 30-130	
m,p-Xylene Column B, Soil	107.819	100.260	100.000000	ND	0.9 107.8	20 30-130	
o-Xylene Column B, Soil	55.7559	60.5420	50.000000	ND	7.3 111.5	20 30-130	
					8.2	20	
LCS Leberatory Control Sample		BXS0621068	157626-1			06/26/2006 072	29
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Methyl tert-Butyl ether, Water Benzene, Water Toluene, Water Ethylbenzene, Water m,p-Xylene, Water o-Xylene, Water Xylenes (total), Water Total BIEX, Water	49.1416 48.6675 49.1170 48.4982 99.1402 49.7578 148.8980 295.1807		50.000000 50.000000 50.000000 50.000000 100.000000 50.000000 150.000000 50.000000		98.3 97.3 98.2 97.0 99.1 99.5 99.3 98.4 97.7	76-123 72-134 76-131 75-131 75-130 74-129 70-130 76-123	
Tert-Butyl Methyl Ether Column B, Water Benzene Column B, Water	48.8600 47.5087		50.000000		95.0	76-123 72-134	

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* %=% REC, R=RPD, A=ABS Diff., D=% Diff.



	Job Number.: 318054	QUALITY	CONTROL	RESULT		Date.: 07/10	0/2006
CUSTOMER: AT	C Associates, Inc.	PROJEC	T: 9003 N. MAI	.	ATTN:		
QC Type	Description		Reag. Code	Lab I	ID Dilut	ion Factor	Date Time
LCS	Laboratory Control Sample		BX\$062106B	157626-1			06/26/2006 0729
Param	eter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	t * Limits F
Toluene Column Ethylbenzene C m,p-Xylene Col o-Xylene Colum	olumn B, Water umn B, Water	48.5180 47.7200 98.3282 48.3311		50.000000 50.000000 100.000000 50.000000		97.0 95.4 98.3 96.7	76-131 75-131 75-130 74-129
MB	Method Blank			157626-1			06/26/2006 0749
Param	eter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	t * Limits F
Benzene, Water Toluene, Water Ethylbenzene, m,p-Xylene, Water Xylenes (total Total BTEX, Water Tert-Butyl Met Benzene Column Toluene Column	Water iter ir), Water iter hyl Ether Column B, Water i B, Water i B, Water olumn B, Water umn B, Water	ND ND ND ND O.0000 ND ND ND ND ND ND ND ND ND ND ND ND ND					
МВ	Method Blank			157626-1	20.0	00	06/26/2006 0819
Param	meter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Resul	t * Limits F
Benzene, SPLP Toluene, SPLP Ethylbenzene, m,p-Xylene, SF o-Xylene, SPLF Xylenes (total Total BTEX, SF	PLP), SPLP thyl Ether Column B, SPLP n B, SPLP n B, SPLP column B, SPLP .umn B, SPLP	ND ND ND ND ND ND O.0000 O.0000 ND ND ND ND ND ND ND ND ND ND					
MS	Matrix Spike		BX\$062306B	317399-1	20.0	00	D6/26/2006 D940
Param	neter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Resul	
Methyl tert-Bu Benzene, SPLP Toluene, SPLP Ethylbenzene,	utyl ether, SPLP	50.4870 43.9578 43.9321 43.0510		50.000000 50.000000 50.000000 50.000000	ND ND ND	88 88 86	70-130 70-130 70-130 70-130
			Page 37	* %=% REC,	R=RPD, A=ABS	Diff., D=% D	iff.



	Job Number.: 318054	QUALITY	CONTROL	RESULT		Date.: 07/10	/2006	
CUSTOMER: AT	C Associates, Inc.	PROJE	CT: 9003 N. MAI	N	ATTN:			
QC Type	Description		Reag. Code	Lab	ID Dilut	ion Factor	Date Time	e
MS	Matrix Spike		BXS062306B	317399-1	20.00	0	06/26/2006 094	40
Param	neter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
m,p-Xylene, SP o-Xylene, SPLP Xylenes (total Total BTEX, SP Tert-Butyl Met Benzene Column Toluene Column Ethylbenzene C m,p-Xylene Colum), SPLP LP thyl Ether Column B, SPLP B B, SPLP B B, SPLP Column B, SPLP cumn B, SPLP	89.7171 44.1903 133.9074 264.8483 50.6467 42.8537 42.8234 42.4284 87.2334 42.8149		100.000000 50.000000 150.000000 50.000000 50.000000 50.000000 50.000000 100.000000	ND 0.0000 0.0000 1.78440 ND ND ND	90 88 89 88 98 86 86 85 87	70-130 70-130 70-130 70-130 70-130 70-130 70-130 70-130 70-130	
MSD	Matrix Spike Duplicate		BXS062306B	317399-1	20,00	0	06/26/2006 100	00
Param	neter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Methÿl tert-Bu	ityl ether, SPLP	52.0804	50.4870	50.000000	1.37053	101.4 3.2	70-130 20	
Benzene, SPLP		44.1746	43.9578	50.000000	ND	88.3 0.5	70-130 20	
Toluene, SPLP		43.9849	43.9321	50.000000	ND	88.0	70-130	
Ethylbenzene,	SPLP	43.3784	43.0510	50.000000	ND	0.1 86.8	20 70-130	
m,p-Xylene, SP	PLP	90.1354	89.7171	100.000000	ND	0.8 90.1	20 70-130	
o-Xylene, SPLP	•	44.5674	44.1903	50.000000	ND	0.5 89.1	20 70-130	
Xylenes (total	.), SPLP	134.7028	133.9074	150.000000	0.0000	0.8 89.8	20 70-130	
Total BTEX, SP	•	266.2407	264.8483	300.000000	0.0000	0.6 88.7	20 70-130	
•	hyl Ether Column B, SPLP	52.1440	50.6467	50.000000	1.78440		20 70-130	
Benzene Column		42.8282	42.8537	50.000000	ND	3.0 85.7	20 70-130	
Toluene Column	•	42.7408	42.8234	50.000000	ND	0.1 85.5	20 70-130	
Ethylbenzene C	•	42.3851	42.4284	50.000000	ND	0.2 84.8	20 70-130	
m,p-Xylene Col	·	87,2536	87.2334	100.000000		0.1 87.3	20 70-130	
o-Xylene Colum	-	42.9718	42.8149	50.000000		0.0 85.9 0.4	20 70-130 20	



							•	
	Job Number.: 318054	QUALITY	CONTROL	. RESULT		t Date.: 07/10/	/2006	
CUSTOMER: A	TC Associates, Inc.	PROJEC	CT: 9003 N. MAI	N	ATTN:			
QC Type	Description		Reag. Code	e Lab	ID Dilu	tion Factor	Date Ti	me
	: TNRCC 1005 ription.: Direct Analytical	TPH Method TX 100		: mg/		Analyst	: тер	
LCD	Laboratory Control Sample	Dupticate	GCL051906	157296-1			06/21/2006 2	012
Parar	meter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hydi	rocarbons C6 - C12, Soil	236.881	234.799	250.000000	ND	95	70-130	
Petroleum Hyd	rocarbons C12 - C28, Soil	275.306	281.051	250.000000	ND	0.9 110	20 70-130	
Petroleum Hydi	rocarbons C6 - C35, Soil	512.187	515.849	500.000000	ND	2.1 102 0.7	20 70-130 20	
LCS	Laboratory Control Sample		GCL051906	157296-1			36/21/2006 1	938
Para	meter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hyd	rocarbons C6 - C12, Soil rocarbons C12 - C28, Soil rocarbons C6 - C35, Soil	234.799 281.051 515.849	•	250.000000 250.000000 500.000000	ND ND ND	93.9 112.4 103.2	70-130 70-130 70-130	
МВ	Method Blenk		GC061306	157296-1		1	06/21/2006 1	905
Para	meter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hyd Petroleum Hyd	rocarbons C6 - C12, Soil rocarbons C12 - C28, Soil rocarbons C28 - C35, Soil rocarbons C6 - C35, Soil	ND ND ND ND						
мѕ	Matrix Spike		GCS061906	318024-2		ı	06/21/2006 2	151
Para	meter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hyd	rocarbons C6 - C12, Soil rocarbons C12 - C28, Soil rocarbons C6 - C35, Soil	232.221 174.600 406.821		250.000000 250.000000 500.000000	ND 173.813 354.574	93 0 10	70-130 70-130 70-130	A A
MSD	Matrix Spike Duplicate		GCS061906	318024-2		ſ	06/21/2006 2	224
Para	meter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hyd	rocarbons C6 - C12, Soil	222.920	232.221	250.000000	ND	89	70-130	
Petroleum Hyd	rocarbons C12 - C28, Soil	184.975	174.600	250,000000	173.813	4.1	20.0 70-130	A
Petroleum Hyd	rocarbons C6 - C35, Soil	407.896	406.821	500.000000	354.574	5.8 11 0.3	20.0 70-130 20.0	A

Page 39 * %=% REC, R=RPD, A=ABS Diff., D=% Diff.



Job Number.: 318054	QUALITY	CONTROL	RESULT		t Date.: 07/10,	/2006	
CUSTOMER: ATC Associates, Inc.	PROJE	CT: 9003 N. MAI	N	ATTN:			
QC Type Description		Reag. Code	Lab	ID Dilu	tion Factor	Date Ti	me
LCD Laboratory Control Sample	Duplicate	GCL 051906	157449-1			06/23/2006 1:	516
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hydrocarbons C6 - C12, Water	337.174	356.667	333.333333	ND	101	70-130	
Petroleum Hydrocarbons C12 - C28, Water	300.856	321.956	333.333333	ND	5.6 90	20 70-130	
Petroleum Hydrocarbons C6 - C35, Water	638.030	678.622	666.666667	ND	6.8 96 6.2	20 70-130 20	
LCS Laboratory Control Sample		GCL051906	157449-1			06/23/2006 1	443
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hydrocarbons C6 - C12, Water Petroleum Hydrocarbons C12 - C28, Water Petroleum Hydrocarbons C6 - C35, Water	356.667 321.956 678.622		333.333333 333.333333 666.666667	ND	107.0 96.6 101.8	70-130 70-130 70-130	
MB Method Blank		GC961306	157449-1		1	06/23/2006 1	410
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hydrocarbons C6 - C12, Water Petroleum Hydrocarbons C12 - C28, Water Petroleum Hydrocarbons C28 - C35, Water Petroleum Hydrocarbons C6 - C35, Water	ND ND ND ND						
SB Spiked Blank		GCS061906	157449-1		(06/23/2006 1	549
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hydrocarbons C6 - C12, Water Petroleum Hydrocarbons C12 - C28, Water Petroleum Hydrocarbons C6 - C35, Water	369.036 387.017 756.053		333.333333 333.333333 666.666667		110.7 116.1 113.4	70-130 70-130 70-130	
SBD Spiked Blank Duplicate		GCS061906	157449-1			06/23/2006 1	623
Parameter/Test Description	QC Result	QC Result	True Value	Orig. Value	Calc. Result	* Limits	F
Petroleum Hydrocarbons C6 - C12, Water	345.353	369.036	333.333333	ND	103.6	70-130 20	
Petroleum Hydrocarbons C12 - C28, Water	316.809	387.017	333.333333	ND	6.6 95.0	70-130	
Petroleum Hydrocarbons C6 - C35, Water	662.162	756,053	666.666667	ND	20.0 99.3 13.2	20 70-130 20	



dor	Number.: 318054	QUALITY	CONTRO	L RESUL		ort Date.: 07/10	/2006	
CUSTOMER: ATC AS	sociates, Inc.	PROJEC	CT: 9003 N. MA	iIN	ATTI	l:		
QC Type	Description		Reag. Coo	le Lab	ID Dil	ution Factor	Date Ti	те
Test Method Method Descripti	: SW-846 8315 on.: Formaldehyde by HF	PLC		: 157630	g/L	Analyst	: jps	
LCS Lab	oratory Control Sample		LC050106	157516			06/26/2006 1	024
Parameter	/Test Description	QC Result	QC Result	True Value	Orig. Value	e Calc. Result	* Limits	F
Formaldehyde, Wate	r	1069.64		10000.		107.0	39-153	
	hod Blank			157516		L		009
Parameter Formaldehyde, Wate	/Test Description r	QC Result	QC Result	True Value	Orig. Value	e Calc. Result	* Limits	-
MS Mat	rix Spike		LC050206	318178-	1		06/26/2006 1	055
Parameter	/Test Description	QC Result	QC Result	True Value	Orig. Value	e Calc. Result	* Limits	F
Formaldehyde, Liqu	iid	991.05		10000.	29.92	96	39-153	
MSD Mat	rix Spike Duplicate		LC050206	318178-	1		06/26/2006 1	110
Parameter	/Test Description	QC Result	QC Result	True Value	Orig. Valu	e Calc. Result	* Limits	F
Formaldehyde, Liqu	ıid	904.06	991.05	10000.	29.92	87.4 9.5	70-130 20	



Method: GC Volatile Organ	cs	Metho	d Code	.: 8021		Prep Batch:
Batch(s): 157626		Test	Equipment Code: BTEX02			
ab ID DT Sample ID		Date	ATFT	ATFTB	BFB	BFBB
7626- 1 LCS 7626- 1 MB 8054- 4 B-1 8054- 5 B-2 8054- 6 B-3		06/26/2006 06/26/2006 06/26/2006 06/26/2006 06/26/2006	107.2 116.0 185.8A 117.5 117.0	108.0 115.1 142.2A 116.0 117.0	105.3 111.9 107.7 113.7 112.2	107.7 113.7 98.2 111.6 116.5
est Test Description	Limits	;				
FT a,a,a-Trifluorotoluene FTB a,a,a-Trifluorotoluene Column B BFB (Surrogate) BB BFB (Surrogate) Column B	70 - 13 B 70 - 13 64 - 13 64 - 13	5 6				
Method: GC Volatile Organ Batch(s): 157626	ics		d Code Matrix			Prep Batch: Equipment Code: BTEX02
ab ID DT Sample ID		Date	ATFT	ATFTB	BFB	BFBB
7626- 1 MB 7399- 1 MS BA1R 4' 7399- 1 MSD BA1R 4'		06/26/2006 06/26/2006 06/26/2006	119.7 114.8 114.8	115.4 115.2 115.0	111.6 109.7 111.0	114.0 111.6 112.3
est Test Description	Limits	.				
FT a,a,a-Trifluorotoluene FTB a,a,a-Trifluorotoluene Column B BFB (Surrogate) BB BFB (Surrogate) Column B	70 - 13 B 70 - 13 64 - 13	5 6				
Method: GC Volatile Organ Batch(s): 157581	ics		d Code Matrix			Prep Batch: Equipment Code: BTEX02
ab ID DT Sample ID		Date	ATFT	ATFTB	BFB	BFBB
7581- 1 LCS 7581- 1 MB 7581- 1 SB 7581- 1 SBD 8054- 1 B-1 12-14 8054- 2 B-2 20-22 8054- 3 B-3 20-22		06/23/2006 06/23/2006 06/23/2006 06/23/2006 06/23/2006 06/23/2006 06/23/2006	104.0 111.3 103.6 106.9 94.6 153.3A 86.1	104.1 113.3 106.8 108.7 96.3 183.1A 86.0	107.2 106.2 93.8 110.7 92.0 15.7A 85.0	102.3 106.0 103.4 106.9 97.5 103.4 91.3
est Test Description	Limits	1				
FT a,a,a-Trifluorotoluene FTB a,a,a-Trifluorotoluene Column B BFB (Surrogate) BB BFB (Surrogate) Column B	50 - 15 B 50 - 15 50 - 15 50 - 15	i0 i0				



SURROGATE RECOVERIES REPORT

Job Number.: 318054

Report Date.: 07/10/2006

CUSTOMER: ATC Associates, Inc.

PROJECT: 9003 N. MAIN

ATTN: Patrick Dworsczyk

Method: Direct Analytical TPH Method TX 1005 Batch(s): 157402 157465				d Code: TX1005 Matrix: Soil	Prep Batch: Equipment Code:			
Lab ID		DT	Sample ID	_	Date	OTERPH		
157296-	1 LC				06/21/2006	93.34		
157296-	1 LC	S			06/21/2006	94.02		
157296-	1 MB				06/21/2006	89.50		
318024-	2 MS		EPO-45-1-(5'-6')		06/21/2006	85.43		
318024-	2 MS)	EPO-45-1-(51-61)		06/21/2006	89.00		
318054-	1		B-1 12-14		06/22/2006	95.08		
318054-	2		8-2 20-22		06/22/2006	93.28		
318054-	3		8-3 20-22		06/22/2006	91.91		
Test	Te	Test Description Limit		Limits				
OTERPH	0-	o-Terphenyl 70 - 13			0			

Method: Direct Analytical TPH Method TX 1005 Batch(s): 157591					ch: 157449 Code: EXTGC12
Lab ID		DT	Sample ID	Date OTERPH	
157449-	1 LCD			06/23/2006 96.46	
157449-	1 LCS			06/23/2006 102.3	
157449-	1 MB			06/23/2006 101.4	
	1 SB			06/23/2006 116.6	
	1 SBD			06/23/2006 104.9	
318054-	4		B-1	06/23/2006 102.8	
318054-	-		8-2	06/23/2006 101.0	
318054-			B-3	06/24/2006 107.3	
Test	Test Description Limit		cription	Limits	
OTERPH	0-Te	erphe	nyl	70 - 130	



QUALITY ASSURANCE METHODS

REFERENCES AND NOTES

Report Date: 07/10/2006

REPORT COMMENTS

- All pages of this report are integral parts of the analytical data. Therefore, this report should be reproduced only in its entirety.
- 2) Reporting limits are adjusted for sample size used, dilutions and moisture content if applicable.
- 3) According to 40CFR Part 136.3, pH, Chlorine Residual, and Dissolved Oxygen analyses are to be performed immediately after aqueous sample collection. When these parameters are not indicated as field,(e.g. pH field) they were not analyzed immediately, but as soon as possible on laboratory receipt.
- 4) For all USACE projects, the QC limits are based on "mean +/- 2 sigma", which are the warning limits.

General Information:

- Cresylic Acid is the combination of o,m and p-Cresol. The combination is reported as the final result.
- m-Cresol and p-Cresol co-elute. The result of the two is reported as either m&p-cresol or as p-cresol.
- m-Xylene and p-Xylene co-elute. The result of the two is reported as m,p-Xylene.
- N-Nitrosodiphenylamine decomposes in the gas chromatograph inlet forming dipheylamine and, consequently, may be detected as diphenylamine.
- Methylene Chloride and Acetone are recognized potential laboratory contaminants. Its presence in the sample up to five times the amount reported in the blank may be attributed to laboratory contamination.
- Trimethysilyl(Diazomethane) is used to esterify acid herbicides in Method SW-846 8151A.
- For Inorganic analyses, duplicate QC limits are determined as follows: If the sample result is less than or equal to 5 times the reporting limit, the RPD limit is equal to the reporting limit. If the sample result is greater than 5 times the reporting limit, the RPD limit is the method defined RPD.
- For TRRP reports, the header on the column RL is equivalent to a MQL/PQL.

Explanation of Qualifiers:

- U This qualifier indicates that the analyte was analyzed but not detected.
- J (Organics only) This qualifier indicates that the analyte is an estimated value between the RL and the MDL.
- B (Inorganics only) This Qualifier indicates that the analyte is an estimated value between the RL and the MDL.
- N (Organics only) This flag indicates presumptive evidence of a compound. This flag is only used for tentatively identified compounds (TICs), where the identification is based on a mass spectral library search. It is applied to all TIC results. For generic charachterization of a TIC, such as "chlorinated hydrocarbon", the "N" flag is not used.

Explanation of General QC Outliers:

- A Matrix interference present in sample.
- a MS/MSD analyses yielded comparable poor recoveries, indicating a possible matrix interference. Method performance is demonstrated by acceptable LCS recoveries.
- b Target analyte was found in the method blank.
- M QC sample analysis yielded recoveries outside QC acceptance criteria. This sample was reanalyzed.
- L LCS analysis yielded high recoveries, indicating a potential high bias. No target analytes were observed above the RL in the associated samples.
- G Marginal outlier within 1% of acceptance criteria.
- r RPD value is outside method acceptance criteria.
- C Poor RPD values observed due to the non-homogenous nature of the sample.
- O Sample required dilution due to matrix interference.
- D Sample reported from a dilution.
- d Spike and/or surrogate diluted.
- P The recovery of this analyte is outside default QC limits. The data is accepted and will be used to calculate in-house statistical limits.
- E The reported concentration exceeds the instrument calibration.
- F The analyte is outside QC limits. The sample data is accepted since this analyte is not reported in associated samples.
- H Continuing Calibration Verification (CCV) standard is not associated with the samples reported.



QUALITY ASSURANCE METHODS REFERENCES AND NOTES

Report Date: 07/10/2006

q - See the subcontract final report for qualifier explanation.

- W The MS/MSD recoveries are outside QC acceptance criteria because the amount spiked is much less than the amount found in the sample.
- K High recovery will not affect the quality of reported results.

Z - See case narrative.

Explanation of Organic QC Outliers:

- e Method blank analysis yielded phthalate concentrations above the RL. Phthlates are recognized potential laboratory contaminants. Its presence in the sample up to five times the amount reported in the blank may be attributed to laboratory contamination.
- S Sample reanalyzed/reextracted due to poor surrogate recovery. Reanalysis confirmed original analysis indicating a possible matrix interference.

T - Sample analysis yielded poor surrogate recovery.

- R The RPD between the two GC columns is greater than 40% and no anomalies are present. The higher result is reported as per EPA Method 8000B.
- I The RPD between the two GC columns is greater than 40% and anomalies are present. The lower of the two results has been reported.

X - Gaseous compound. In-house QC limits are advisory.

- Ketone compounds have poor purge efficiency. In-house QC limits are advisory.

f - Surrogate not associated with reported analytes.

Explanation of Inorganic QC Outliers:

- Q Method blank analysis yielded target analytes above the RL. Associated sample results are greater than 10 times the concentrations observed in the method blank.
- The RPD control limit for sample results less than 5 times the RL is +/- the RL value. Sample and duplicate results are within method acceptance criteria.

e - Serial dilution failed due to matrix interference.

- g Sample result quantitated by Method of Standard Additions (MSA) due to the analytical spike recovery being below 85 percent. The correlation coefficent for the MSA is greater than or equal to 0.995.
- BOD/cBOD seed value is not within method acceptance criteria. Due to the nature of the test method, the sample cannot be reanalyzed.
- 1 BOD/CBOD LCS value is not within method acceptance criteria. Due to the nature of the test method, sample cannot be reanalyzed.

- Spiked sample recovery is not within control limits.

n - Sample result quantitated by Method of Standard Additions (MSA) due to the analytical spike recovery being below 85 percent. The correlation coefficient for the MSA is less than 0.995.

* - Duplicate analysis is not within control limits.

Abbreviations:

- Designation given to identify a specific extraction, digestion, preparation, or analysis set. Batch

- Continuing Calibration Verification CCV

- Low level standard check - GFAA, Mercury CRA

- Low level standard check - ICP CR I

Dil Fac - Dilution Factor - Secondary dilution analysis

- Detection Limit Factor DLFac

- Duplicate DU

- Extraction Blank (TCLP, SPLP, etc.)

EB - Initial Calibration ICAL

ICB - Initial Calibration Blank

ICV Initial Calibration Verification

Interference Check Sample A - ICP ISA ISR

- Interference Check Sample B - ICP - Laboratory Control Duplicate LCD

LCS - Laboratory Control Sample



QUALITY ASSURANCE METHODS REFERENCES AND NOTES

Report Date: 07/10/2006

MB - Method Blank - Method Duplicate MD - Method Detection Limit MDI - Method Quantitation Limit (TRRP) MQL - Matrix Spike - Matrix Spike Duplicate MSD - Not Detected ND - Preparation Blank PR PREPF - Preparation Factor - Reporting Limit RPD - Relative Percent Difference - Relative Response Factor RRF - Retention Time RT - Sample Quantitation Limit (TRRP) SQL TIC - Tentatively Identified Compound

Method References:

- (1) EPA 600/4-79-020 Methods for the Analysis of Water and Wastes, March 1983.
- (2) EPA 600/R-94-111 Methods for the Determination of MEtals in Environmental Samples, Supplement I, May 1994.
- (3) EPA SW846 Test Methods for Evaluating Solid Waste, Third Edition, September 1986; Update I July 1992; Update II, September 1994, Update IIA August 1993; Update IIB, January 1995; Update III, December 1996, Update IVA January 1998, Update IVB November 2000.
- (4) Standard Methods for the Examination of Water and Wastewater, 16th Edition (1985), 17th Edition (1989), 18th Edition (1992), 19th Edition (1995), 20th Edition (1998). (5) HACH Water Analysis Handbook 3rd Edition (1997).

- Federal Register, July 1, 1990 (40 CFR Part 136 Appendix A).

 Compendium of Methods for the Determination of Toxic Organic Compounds in Ambient Air, 2nd Edition, (7) January 1997.
- (9) Diagnosis and Improvement of Saline and Alkali Soils, Agriculture Handbook No. 60, United States Department of Agriculture, 1954.



LABORATORY CHRONICLE

Job Number: 318054

Date: 07/10/2006

CUSTOMER: ATC Ass	ociates, inc. PROJE	CT: 9003 N	. MAIN		A	TTN: Patrick D	lworaczy	k
ab ID: 318054-1	Client ID: B-1 12-14	Date Re	cvd: 06/	21/2006	Sample	Date: 06/21/20	006	
METHOD	DESCRIPTION			PREP BT		DATE/TIME AN		DILUTIO
METHOD	Data Package Validation	1	158535		"(")	07/10/2006	0000	0.20.70
THROC 100E	Direct Analytical TPH Method TX 1005	i		157296		06/22/2006	2038	1.0000
TNRCC 1005	Direct Analytical Irm Method IA 1005	i	157594	131270		06/26/2006	1045	,,,,,,,,
	Extractable GC Data Package Production						0800	
SW846 8315	Extraction Formaldehyde Waters	1	157516	457547		06/23/2006		4 000
sw-846 8315	Formaldehyde by HPLC	1		157516		06/26/2006	1156	1.000
SW-846 8021B	GC Volatile Organics	1	157581			06/23/2006	2050	1.0000
	GC Volatiles Data Package Production	1						
SM-2540 G Mod.	Moisture (Total + Fixed Solids, Ash)	1	157440			06/22/2006	1630	
TNRCC TX-1005	TNRCC 1005 Extraction (Ultrasonic)	1	157296			06/21/2006	1600	
SW-846 1311	Toxicity Characteristic Leachate Proced.	1	157473			06/22/2006	2230	
ab ID: 318054-2	Client ID: B-2 20-22	Date Re	cvd: 06/	21/2006	Sample	Date: 06/21/20	006	
METHOD	DESCRIPTION			PREP BT		DATE/TIME AN		DILUTIO
	Direct Analytical TPH Method TX 1005	1		157296	W(U)	06/22/2006	2144	1.0000
TNRCC 1005				131290				1.0000
	Extractable GC Data Package Production	1	157594			06/26/2006	1045	
SW846 8315	Extraction Formaldehyde Waters	1	157516			06/23/2006	0800	
SW-846 8315	Formaldehyde by HPLC	1		157516		06/26/2006	1212	1.000
SW-846 80218	GC Volatile Organics	1	157581			06/23/2006	2130	1.0000
SM-2540 G Mod.		1	157440			06/22/2006	1630	
TNRCC TX-1005	TNRCC 1005 Extraction (Ultrasonic)	1	157296			06/21/2006	1600	
SW-846 1311	Toxicity Characteristic Leachate Proced.	i	157473			06/22/2006	2230	
		D	مرساء ٥٤ ١	734 /2004	Formia	Data 06/21/2	204	
ab ID: 318054-3	Client ID: B-3 20-22			21/2006		Date: 06/21/20		
METHOD	DESCRIPTION			PREP BT	#(S)	DATE/TIME A		DILUTI
TNRCC 1005	Direct Analytical TPH Method TX 1005	1		157296		06/22/2006	2217	1.0000
	Extractable GC Data Package Production	1	157594			06/26/2006	1045	
SW846 8315	Extraction Formaldehyde Waters	1	157516			06/23/2006	0800	
SW-846 8315	Formaldehyde by HPLC	1	157630	157516		06/26/2006	1227	1.000
SW-846 8021B	GC Volatile Organics	1	157581			06/23/2006	2230	1.0000
		i	157440			06/22/2006	1630	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
SM-2540 G Mod.		j	157296				1600	
TNRCC TX-1005	TNRCC 1005 Extraction (Ultrasonic)	•				06/21/2006		
sw-846 1311	Toxicity Characteristic Leachate Proced.	1	157473			06/22/2006	2230	
ab ID: 318054-4	Client ID: B-1		cvd: 06/			Date: 06/21/20		
METHOD	DESCRIPTION			PREP BT	#(S)	DATE/TIME AN		DILUTIO
TNRCC 1005	Direct Analytical TPH Method TX 1005	1	157591	157449		06/23/2006	2303	1.0000
SW846 8315	Extraction Formaldehyde Waters	1	157516			06/23/2006	0800	
SW-846 8315	Formaldehyde by HPLC	1	157630	157516		06/26/2006	1258	1.000
SW-846 8021B	GC Volatile Organics	1	157626			06/26/2006	1020	1.0000
TNRCC TX-1005	TX-1005 Extraction Water	i	157449			06/22/2006	1430	,,,,,,,,,
-1- rh - 7400F/ F	nii In- n-3	Doto Do	cvd: 06/	21 /2006	Comple	Date: 04/21/20	204	
.ab ID: 318054-5	Client ID: B-2					Date: 06/21/20		DYLLET
METHOD	DESCRIPTION			PREP BT	#(5)	DATE/TIME AN		DILUTIO
TNRCC 1005	Direct Analytical TPH Method TX 1005	1		157449		06/23/2006	2336	1.0000
SW846 8315	Extraction Formaldehyde Waters	1	157516			06/23/2006	0800	
SW-846 8315	Formaldehyde by HPLC	1	157630	157516		06/26/2006	1313	1.000
SW-846 80218	GC Volatile Organics	1	157626			06/26/2006	1040	1.0000
TNRCC TX-1005	TX-1005 Extraction Water	1	157449			06/22/2006	1430	
ab ID: 318054-6	Client ID: B-3	Date Pe	cvd: 06/	21/2006	Sample	Date: 06/21/20	006	
								DILUTI
	DESCRIPTION	_		PREP BT	#(0)	DATE/TIME AN		
METHOD	Direct Analytical TPH Method TX 1005	1	12/271	157449		06/24/2006	0009	1.0000
TNRCC 1005								
	Extraction Formaldehyde Waters	1	157516			06/23/2006	0800	
TNRCC 1005		1		157516		06/23/2006 06/26/2006	0800 1329	1.000



LABORATORY CHRONICLE

Job Number: 318054

Date: 07/10/2006

CUSTOMER: ATC Associates, Inc.

PROJECT: 9003 N. MAIN

ATTN: Patrick Dworaczyk

Lab ID: 318054-6 METHOD

Client ID: B-3
DESCRIPTION

TNRCC TX-1005 TX-1005 Extraction Water

Date Recvd: 06/21/2006 Sample Date: 06/21/2006

RUN# BATCH# PREP BT #(S) 157449

DATE/TIME ANALYZED DILUTION

06/22/2006 1430